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(54) Title: ANTIGENIC PEPTIDES, SUCH AS FOR G PROTEIN-COUPLED RECEPTORS (GPCRS), ANTIBODIES THERETO, AND SYSTEMS FOR IDENTIFYING SUCH ANTIGENIC PEPTIDES

(57) Abstract: The present invention provides antigenic peptides for GPCRs and antibodies relating thereto, and related systems, methods, compositions, and the like, such as diagnostics and medicaments. Where antibodies against a given GPCR are not known, the present invention provides such antibodies, and preferred antigenic sequences for producing such antibodies. Where antibodies against a given GPCR are known, the present invention provides preferred antigenic peptides for producing antibodies that exhibit improved specificity, affinity or capacity to perform antibody-related actions relative to the known antibodies.

ANTIGENIC PEPTIDES, SUCH AS FOR G PROTEIN-COUPLED RECEPTORS (GPCRS), ANTIBODIES THERETO, AND SYSTEMS FOR IDENTIFYING SUCH ANTIGENIC PEPTIDES

5 CROSS-REFERENCE TO RELATED APPLICATIONS

[1] The present application claims priority from United States provisional patent application No. 60/257,144, filed December 19, 2000 and presently pending.

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- [2] The following is a Table of Contents to assist review of the present application:
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ANTIGENIC PEPTIDES GENERALLY:

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 - E. CERTAIN ASSAYS, ANTIBODIES, PROBES, THERAPEUTICS, AND OTHER SYSTEMS AND ASPECTS, OF THE INVENTION
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SCREENING FOR/WITH ANTIGENIC PEPTIDES:

LIST OF ASSAYS:

ENZYME-LINKED IMMUNOSORBENT ASSAYS (ELISA):

IMMUNOFLUORESCENCE ASSAY:

BEAD AGGLUTINATION ASSAYS:

ENZYME IMMUNOASSAYS:

SANDWICH ASSAY:

SEQUENTIAL AND SIMULTANEOUS ASSAYS:

IMMUNOSTICK (DIP-STICK) ASSAYS:

IMMUNOCHROMATOGRAPHIC ASSAYS:

IMMUNOFILTRATION ASSAYS:

BIOSENSOR ASSAYS:

ANTIBODIES ANTIBODIES GENERATED AGAINST A PARTICULAR ANTIGENIC PEPTIDE AND ITS CORRESPONDING GPCR: ANTIBODIES GENERALLY: ANTI-IDIOTYPIC ANTIBODIES: a. Antibody Preparation Polyclonal Antibodies (i) ANTIBODY PREP - POLYCLONAL: ANTIBODY PREP - ADJUVANTS (ALL ABS): Monoclonal Antibodies 10 (ii) ANTIBODY PREP - MONOCLONAL: MOABS - COMBINATORIAL: **HUMANIZED MOAB:** ANTIBODY SUBSTITUTIONS - NON-IMMUNOGLOBULIN POLYPEPTIDES 15 (ALL ABS): CHIMERICS: ANTIBODY LABELING (ALL ABS): (iii) Humanized And Human Antibodies **HUMANIZED AB GENERALLY:** 20 (iv) Antibody Fragments ANTIBODY FRAGMENTS: (v) Bispecific Antibodies **BISPECIFIC ANTIBODIES GENERALLY:** ANTIBODIES - HYBRID IMMUNOGLOBULIN HEAVY CHAIN: ANTIBODIES - CROSS-LINKED OR "HETEROCONJUGATE": 25 ANTIBODIES - DIABODIES: ANTIBODIES - OTHER: Ъ. **Antibody Purification** ANTIBODY PURIFICATION GENERALLY: 30 **BEFORE LPHIC:** LPHIC: POST LPHIC: c. Some Uses For Antibodies Described Herein Generally GENERALLY: 35 ASSAYS: DIAGNOSTIC USES: (ii) Assays ASSAYS: COMPETITIVE BINDING ASSAYS: 40 Affinity Purification (iii) AFFINITY PURIFICATION: Therapeutics (iv) THERAPEUTIC USES: 45 THERAPEUTIC FORMULATIONS: THERAPEUTIC FORMULATIONS -STERILE: THERAPEUTIC ADMINISTRATIONS:

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5. DRUG DESIGN BASED ON THE ANTIGENS HEREIN OR ANTIBODIES THERETO

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BACKGROUND

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- [4] G protein-coupled receptors (GPCRs) are a large group of proteins that transmit signals across cell membranes. In general terms, GPCRs function somewhat like doorbells. When a molecule outside the cell contacts the GPCR (pushes the doorbell), the GPCR changes its shape and activates "G proteins" inside the cell (similar to the doorbell causing the bell to ring inside the house, which in turn causes people inside to answer the door). GPCRs are like high-security doorbells because each GPCR responds to only one specific kind of signaling molecule (called its "endogenous ligand"), kind of like a high-tech door lock that responds to only one fingerprint. Part of the GPCR is located outside the cell (the "extracellular domain"), part spans the cell's membrane (the "transmembrane domain"), and part is located inside the cell (the "intracellular domain"). Thus, GPCRs are embedded in the outer membrane of a cell and recognize and bind certain signaling molecules that are present in the spaces surrounding the cell. GPCRs are used by cells to keep an eye on the cells' own activity and on the environment. In organisms that have many cells, the cells use GPCRs to talk to each other.
 - [5] GPCRs are important to the pharmaceutical industry and other industries. For example, many drugs, including some antibody-based drugs, act by binding to specific GPCRs and initiating or inhibiting their intracellular actions, and diagnostics and therapeutics based on GPCRs or on antibodies for GPCRs are becoming increasingly important.
 - [6] General concepts about GPCRs are discussed in more scientific terms in the following paragraphs.
 - [7] The GPCR superfamily has at least 250 members, Strader et al., FASEB J., 9:745-754 (1995); Strader et al., Annu. Rev. Biochem., 63:101-32 (1994). GPCRs play important

roles in diverse cellular processes including cell proliferation and differentiation, leukocyte migration in response to inflammation, gene transcription, vision (the rhodopsins), smell (the olfactory receptors), neurotransmission (muscarinic acetylcholine, dopamine, and adrenergic receptors), and hormonal response (luteinizing hormone and thyroid-stimulating hormone receptors). Strader et al., supra; U.S. Patent nos. 5,994,097 and 6,063,596. Many important drugs produce their therapeutic actions through their interaction with GPCRs.

Nucleotide and amino acid sequences for many GPCRs have been reported and can be found in public databases such as GenBank and GenPept. Generally speaking, different GPCRs show both structural and sequence similarities. The most conserved domains of 10 GPCRs are the transmembrane domains and the first two cytoplasmic loops. GPCRs range in size from under 400 to over 1000 amino acids. Coughlin, S. R., Curr. Opin. Cell Biol. 6:191-. 197 (1994). They contain seven hydrophobic transmembrane regions that span the cellular membrane and form a bundle of antiparallel alpha helices. McKee K.K., supra. The bundle of helices forming the transmembrane regions provide many structural and functional features of the receptor. In most cases, the bundle of helices form a pocket that binds a signaling molecule. However, when the binding site accommodates larger molecules, the extracellular N-terminal segment or one or more of the three extracellular loops participate in binding and in subsequent induction of conformational change in the intracellular portions of the receptor. These helices are joined at their ends by three intracellular and three extracellular loops. GPCRs also contain cysteine disulfide bridges between the second and third extracellular loops, an extracellular N-terminus, and a cytoplasmic or intracellular Cterminus. The N-terminus is often glycosylated, while the C-terminus is generally phosphorylated. A conserved, acidic-Arg-aromatic triplet present in the second cytoplasmic loop may interact with G Proteins. Most GPCRs contain a characteristic consensus pattern. Watson, S. and S. Arkinstall, The G protein Linked Receptor Facts Book, Academic Press, San Diego, CA (1994); Bolander, F. F. Molecular Endocrinology, Academic Press, San Diego, CA (1994).

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Although GPCRs have many features in common, each GPCR has its own unique [9] characteristics as well. GPCRs have varying nucleotide and amino acid sequences, and 30 varying antigenicity. GPCRs bind a diverse array of specific, extracellular signaling molecules (which can also be referred to as "ligands") including peptides, cytokines, hormones, neurotransmitters, growth factors, and specialized stimuli such as photons,

flavorants, and odorants. Identified ligands include, for example, purines, nucleotides (e.g., adenosine, cAMP, NTPs), biogenic amines (e.g., epinephrine, norepinepherine, dopamine, histamine, noradrenaline, serotonin), acetylcholine, peptides (e.g., angiotensin, calcitonin, chemokines, corticotropin releasing factor, galanin, growth hormone releasing hormone, gastric inhibitory peptide, glucagon, neuropeptide Y, neurotensin, opioids, thrombin, secretin, somatostatin, thyrotropin releasing hormone, vasopressin, vasoactive intestinal peptide), lipids and lipid-based compounds (e.g., cannabinoids, platelet activating factor), excitatory and inhibitory amino acids (e.g., glutamate, GABA), ions (e.g., calcium), and toxins.

- [10] In general, a GPCR binds only one type of signaling molecule and GPCRs are classified according to subfamilies based upon their selectivity and specificity for a particular ligand. When the ligand for a receptor is not known, the receptor is known as an orphan receptor. The extracellular domain interacts with or binds to certain signaling molecules or ligands located outside of the cell. The binding of a ligand to the extracellular domain alters the conformation of the receptor's intracellular domain causing the activation of a G protein. The G protein then activates or inactivates a separate plasma-membrane-bound enzyme or ion
- channel. This chain of events alters the concentration of one or more intracellular messengers (second messengers) such as cyclic AMP (cAMP), inositol triphosphate, diacylglycerol, or Ca²⁺. These, in turn, alter the activity of other intracellular proteins such as cAMP-dependent protein kinase and Ca²⁺/calmodulin-dependent protein kinases, leading to the transduction and amplification of the original extracellular signal. Baldwin, J.M., Curr. Opin. Cell Biol. 6:180-190 (1994). The G protein is deactivated by hydrolysis of GTP by GTPase. U.S. Patent Nos. 5,994,097 and 6,063,596.
- [11] GPCR mutations, both of the loss-of-function and of the activating variety, have been associated with numerous human diseases, Coughlin, *supra*. For example, retinitis pigmentosa may arise from either loss-of-function or activating mutations in the rhodopsin gene. Somatic activating mutations in the thyrotropin receptor cause hyperfunctioning thyroid adenomas, Parma, J. et al., Nature 365:649-651 (1993). Parma et al. indicate that it may be possible that certain G protein-coupled receptors susceptible to constitutive activation may behave as proto-oncogenes. Interestingly, GPCRs have functional homologues in human cytomegalovirus and herpesvirus, so GPCRs may have been acquired during evolution for viral pathogenesis, Strader et al., FASEB J., 9:745-754 (1995); Arvanitakis et al., Nature, 385:347-350 (1997); Murphy, Annu. Rev. Immunol. 12:593-633 (1994). The

importance of the GPCR superfamily is further highlighted by the recent discoveries that some of its family members, the chemokine receptors CXCR4/Fusin and CCR5, are coreceptors for T cell-tropic and macrophage-tropic HIV virus strains, respectively, Alkhatib et al., Science, 272:1955 (1996); Choe et al., Cell, 85:1135 (1996); Deng et al., Nature, 381:661 (1996); Doranz et al., Cell, 85:1149 (1996); Dragic et al., Nature, 381:667 (1996); Feng et al., Science, 272:872 (1996). It is conceivable that blocking these receptors may prevent infection by the human immunodeficiency (HIV) virus. Other GPCR-related items include regulating cellular metabolism and diagnosing, treating and preventing particular diseases associated with particular GPCRs.

- [12] One important way to evaluate GPCRs and antibodies for GPCRs as novel drug targets and for other purposes such as diagnostics is through the creation and use of databases. Such databases can provide large amounts of information about genes, proteins, and other biological matter. An excellent example of such a database is the GPCR database created and maintained by LifeSpan BioSciences, Inc., Seattle, Washington, USA, which database is available by subscription to researchers and others needing such information. The information in the databases can, for example, be searched, compared, and analyzed. The compilation of such databases, as well as the searching, comparing, etc., of the databases, can be referred to as the field of "bioinformatics." Investigations largely related to genes, such as the information found from the sequencing of the human genome, can be called "genomics" while similar activities on proteins can be called "proteomics."
- [13] There has gone unmet a need for improved systems, compositions, methods, and the like relating to improved antigenicity of peptides from GPCRs and antibodies relating thereto. The present invention provides these and other advantages.

SUMMARY

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The present invention provides antigenic peptides for GPCRs and antibodies relating thereto, and related systems, methods, compositions, and the like, such as diagnostics and medicaments. Where antibodies against a given GPCR are not known, the present invention provides such antibodies, and preferred antigenic sequences for producing such antibodies. Where antibodies against a given GPCR are known, the present invention provides preferred antigenic peptides for producing antibodies that exhibit improved specificity, affinity or capacity to perform antibody-related actions relative to the known

antibodies. The present invention also provides improved methods of selecting antigenic peptides from any desired protein or polypeptide, as well as antigenic peptides so produced and antibodies against such antigenic peptides.

The antigenic peptides and antibodies herein can be used, for example, to detect the presence or absence of corresponding GPCRs. They can be used to diagnose a variety of diseases and disorders in which GPCRs are involved, such as, e.g., immune-related diseases, cell growth-related diseases, cell regeneration-related diseases, immunological-related cell proliferative diseases, and autoimmune diseases. Examples of specific diseases include AIDS, allergies, Alzheimer's disease, amyotrophic lateral sclerosis, atherosclerosis, bacterial, fungal, protozoan and viral infections, benign prostatic hypertrophy, bone diseases (e.g., osteoarthritis, osteoporosis), carcinoma (e.g., basal cell carcinoma, breast carcinoma, embryonal carcinoma, ovarian carcinoma, renal cell carcinoma, lung adenocarcinoma, lung small cell carcinoma, pancreatic carcinoma, prostate carcinoma, transitional carcinoma of the bladder, squamous cell carcinoma, thyroid carcinoma), cardiomyopathy, chronic and acute inflammation, circadian rhythm disorders, COPD, Crohn's disease, diabetes, Duchenne muscular dystrophy, embryonal carcinoma, endotoxic shock, environmental stress (e.g., by heat, UV or chemicals), gastrointestinal disorders, glioblastoma multiform, graft vs. host disease, Hodgkin's disease, inflammatory bowel disease, ischemia, stroke, lymphoma, macular degeneration, malignant cytokine production, malignant fibrous histiocytoma, melanoma, meningioma, mesothelioma, multiple sclerosis, nasal congestion, pain, Parkinson's disease, prostate carcinoma, psoriasis, rhabdomyosarcoma, psychotic or neurological disorders (e.g., anxiety, depression, schizophrenia, dementia, mental retardation, memory loss, epilepsy, locomotor problems, respiratory disorders, asthma, eating/body weight disorders including obesity, bulimia, diabetes, anorexia, nausea, hypertension, hypotension), renal disorders, reperfusion injury, rheumatoid arthritis, sarcoma (e.g., chondrosarcoma, Ewing's sarcoma, osteosarcoma), septicemia. seminoma, sexual/reproductive disorders, tonsil, transitional carcinoma of the bladder, transplant rejection, trauma, tuberculosis, ulcers, ulcerative colitis, urinary retention, vascular and cardiovascular disorders, or any other disease or disorder in which G protein-coupled receptors are involved, as well as learning and/or memory disorders, diabetes, pain perception disorders, anorexia, obesity, hormonal release problems, or any other disease or disorder in which a specific GPCR is involved.

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[16] The association of particular GPCRs with particular diseases, disorders or conditions will be apparent to a person of ordinary skill in the art in view of the present application, and thus the association with the antibodies of the present invention to the corresponding diseases, disorders or conditions.

- 5 [17] Thus, in one aspect the present invention provides isolated antigenic peptides according to any one of SEQ ID NOS. 692-2292. The isolated antigenic peptides also comprise an amino acid sequences that are at least about 90% or 95% identical to such sequences, or be an analog of such sequences, or comprise a short antigenic amino acid sequence that is identical to at least 5 consecutive amino acids set forth in any one of such sequences or contain no more than one conservative amino acid substitution over at least 7 consecutive amino acids set forth in any of such sequences. The present invention also provides antibodies, particularly isolated antibody having high specificity and high affinity or avidity for a particular GPCR or other target polypeptide or protein, generated using the antigenic peptides discussed herein.
- 15 [18] The present invention also provides isolated nucleic acid molecules encoding an antigenic peptide or antibody as described herein. The molecule can encode a naturally occurring human antigenic peptide. In some embodiments, the present invention provides processes for producing an isolated polynucleotide can comprise hybridizing a nucleotide encoding an antigenic peptide as discussed herein to DNA such as genomic DNA under stringent or highly stringent conditions and isolating the polynucleotide detected with the nucleotide.
 - [19] The present invention also provides kits and assays, such as kits for the detection of antibodies against a particular GPCR or other target polypeptide in a sample comprising: a) an isolated antigenic peptide as discussed herein and derived from the particular GPCR, and b) at least one of a reagent or a device for detecting the antibodies, or comprising: a) an isolated antibody as described herein, and b) at least one of a reagent or a device for detecting the antibody. The assays include detection of a particular GPCR in a sample, comprising: a) providing an isolated antigenic peptide, b) contacting the isolated antigenic peptide corresponding to the particular GPCR with the sample under conditions suitable and for a time sufficient for the antigenic peptide to bind to one or more antibodies specific for the target protein present in the sample, to provide an antibody-bound target protein, and c) detecting the antibody-bound antigenic peptide, and therefrom determining whether the

sample contains the particular GPCR. The assays can further comprise the step of binding the isolated antigenic peptide or the antibody to a solid substrate, and the sample can be an unpurified sample, for example from a human being.

- [20] The assay can be selected from the group consisting of a countefcurrent immuno5 electrophoresis (CIEP) assay, a radioimmunoassay, a radioimmunoprecipitation, an enzymelinked immuno-sorbent assay (ELISA), a dot blot assay, an inhibition or competition assay, a
 sandwich assay, an immunostick (dip-stick) assays, a simultaneous assay, an
 immunochromatographic assay, an immunofiltration assay, a latex bead agglutination assay,
 an immunofluorescent assay, a biosensor assay, and a low-light detection assay.
- In other aspects, the present invention provides methods of identifying an amino acid sequence for an antigenic peptide from a candidate polypeptide sequence such as a polypeptide or protein wherein the antigenic peptide has a length of about 5 to about 100 amino acids, typically 6 amino acids to about 50 amino acids, and preferably 7 amino acids to about 20 amino acids. The methods comprise: a) searching the candidate polypeptide sequence using a comparison window of the length, and b) selecting against amino acid sequences of the length and having at least 1 to 3 or 4 characteristics selected from the group consisting of 1) at least two consecutive prolines, 2) at least two consecutive serines, 3) at least two consecutive aspartic acids, 6) at least two consecutive glutamic acids, 7) methionine, 8) tryptophan, and 9) at least five consecutive amino acids comprising no charged amino acids. Preferably, the method comprises selecting against at least 5 to all of the characteristics.
 - [22] The methods can comprise, independently or in addition, selecting against amino acid sequences of the desired length having at least one of the following characteristics 1) sequences having at least 5 consecutive amino acids that are identical to an alternative amino acid sequence from an alternative polypeptide that can be different from the candidate polypeptide, 2) posttranslational modification sites, and 3) highly hydrophobic sequences. The posttranslational modification sites can be phosphorylation or glycosylation sites. The methods can also comprise performing a BLAST-type or a FAST-type analyses for the candidate polypeptide sequence.
- These and other aspects, features, and embodiments are set forth within this application, including the following Detailed Description and attached drawings. The present invention comprises a variety of aspects, features, and embodiments; such multiple aspects,

features, and embodiments can be combined and permuted in any desired manner. In addition, various references are set forth herein, including in the Cross-Reference To Related Applications, that discuss certain compositions, apparatus, methods, or other information; all such references are incorporated herein by reference in their entirety and for all their teachings and disclosures, regardless of where the references may appear in this application.

BRIEF DESCRIPTION OF THE DRAWING

- [24] Figure 1 depicts representative examples of the nucleotide and amino acid sequences of the GPCRs for which antigenic peptides are set forth herein, SEQ ID NOS. 1 691.
- 10 [25] Figure 2 depicts amino acid sequences for the antigenic peptides for the GPCRs herein, SEQ ID NOS. 692-2292.
 - [26] Figure 3 depicts a listing of GPCRS for which commercially available antibodies are putatively available.

DETAILED DESCRIPTION

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A. INTRODUCTION AND OVERVIEW

- [27] Diseases such as immune-related diseases, cell growth-related diseases, cell regeneration-related diseases, immunological-related cell proliferative diseases, and autoimmune diseases are serious health problems in the modern world. Any improvement in the diagnosis, treatment or other remediation of such diseases is a significant advance for millions of people. The present invention provides methods of identifying and selecting desirable antigenic peptides for GPCRs and other desired target or candidate proteins and polypeptides. The present invention also provides the antigenic peptides themselves, as well as antibodies against the antigenic peptides (and against proteins or polypeptides containing such antigenic peptides), and related diagnostics, antibody-based therapeutics directed to certain diseases and conditions, and other helpful compositions, systems, kits, assays and the like. The compositions, methods, and the like can be useful, for example, as agonists, antagonists, probes, and otherwise as may be desired.
- [28] The antigenic peptides have been carefully selected using specific selection criteria and methodologies set forth herein to take advantage of particularly advantageous regions of the GPCRs from which they have been derived to provide unusually specific and

immunogenic antigens. These antigenic peptides are particularly useful for producing highly specific antibodies against the antigenic peptides, which, in turn, also means antibodies that are highly specific for the corresponding GPCRs containing the antigenic peptides. Accordingly, the antigenic peptides of the present invention, and the antibodies produced therefrom, are particularly useful for high specifity, low noise diagnostics and, in the case of the antibodies, for certain antibody-based therapeutics, as well as methods, kits, systems, and the like incorporating or based on such antigenic peptides or antibodies.

- [29] The antibodies produced using the antigenic peptides of the present invention, for example, have a specificity for the corresponding GPCR such that the antibodies can selectively detect the corresponding GPCR in a sample containing non-desired or contaminating proteins or polypeptides, such as a tissue or blood sample. Preferably, the antibodies have a high specificity such that no significant amounts of such proteins or polypeptides are detected, and further preferably have a specificity such that only insubstantial to essentially zero amounts of non-desirable proteins are detected.
- 15 [30] The antibodies produced using the antigenic peptides of the present invention, for example, typically have an affinity or avidity constant (Ka) of at least about 10⁷ liters/mole, typically a high affinity or avidity at least about 10⁹ liters/mole, preferably at least about 10¹⁰ liters/mole, and further preferably at least about 10¹¹ liters/mole.
- [31] Figure 1 sets forth the DNA and protein sequences for the GPCRs from which the antigenic peptides of the present invention were derived SEQ ID NOS. 1-691. Figure 2 sets forth the amino acid sequences of exemplary antigenic peptides, SEQ ID NOS. 692-2292. The sequences in Figures 1 and 2 are listed according to SEQ ID NO and LSID, which is an identification number assigned to the given sequence in the LifeSpan Biosciences databases. The sequences in Figure 2 also include an identifier LPID, which is also an identification number assigned to the given sequence in the LifeSpan Biosciences databases. Figure 3 depicts GPCRs for which it has been reported that antibodies are commercially available, SEQ ID NOS. 1, 3, 5, 11, 13, 15, 21, 23, 25, 27, 29, 31, 35, 37, 39, 41, 43, 45, 49, 51, 53, 57, 59, 61, 63, 65, 67, 69, 70, 71, 73, 75, 77, 79, 83, 85, 97, 99, 101, 103, 105, 107, 113, 115, 117, 121, 125, 135, 139, 143, 145, 147, 151, 155, 157, 159, 161, 169, 171, 173, 175, 177, 183, 185, 187, 189, 191, 192, 194, 200, 202, 206, 208, 214, 216, 218, 228, 236, 238, 240, 248, 250, 264, 295, 299, 301, 305, 311, 313, 315, 317, 319, 321, 323, 325, 327, 329, 331, 333, 335, 337, 347, 349, 351, 361, 365, 367, 369, 371, 377, 379, 385, 387, 389, 391, 397,

423, 435, 439, 457, 459, 461, 462, 468, 470, 472, 503, 507, 515, 535, 537, 546, 548, 552, 562, 628, 636; Applicants do not represent that any of the antibodies in Figure 3 that such antibodies are actually commercially available nor that they have any significant specificity nor affinity for the GPCRs reported. For GPCRs for which no antigens or antibodies were previously known, the present invention provides valuable antigenic peptides and antibodies (see, e.g., SEQ ID NOS. 704-712, 731-743, 774-777, 803-806, 821-824, 876-879, 890-916, 942-949, 965-970, 985-988, 994-1009, 1014-1020, 1025-1028, 1044-1048, 1053-1056, 1073-1086, 1114-1123, 1152-1160, 1173-1178, 1188-1197, 1210-1227, 1232-1244, 1258-1270, 1280-1303, 1309-1368, 1373-1377, 1386-1389, 1394-1402, 1462-1482, 1496-1525, 1542-1549, 1557-1563, 1583-1649, 1656-1679, 1684-1688, 1693-1732, 1744-1752, 1765-1839, 1846-1854, 1855-1866, 1871-1917, 1926-1941, 1952-1955, 1960-1980, 1985-2141, 2152-2165, and 2170-2292.); for GPCRs for which antigens or antibodies are known, the present invention provides improved antigens in the form of antigenic peptides and improved antibodies (see, e.g., SEQ ID NOS. 692-703, 713-730, 744-802, 807-820, 825-875, 880-889, 917-941, 950-964, 971-984, 989-993, 1010-1013, 1021-1024, 1029-1043, 1049-1052, 1057-1072, 1087-1113, 1124-1151, 1161-1172, 1179-1187, 1198-1209, 1228-1231, 1245-1257, 1271-1279, 1304-1308, 1369-1372, which are antigenic peptides derived from GPCRs for which antibodies are reportedly commercially available). The antigenic peptides and antibodies, and uses and assays, etc., related to the antigenic peptides, are discussed further below.

[32] The discussion herein, including the following passages, has been separated by headings for convenience. The disclosure under a given heading is not restricted to that heading. For example, the discussion in the definitions section is a part of the disclosure of the invention, the discussion on antigenic peptides also contains discussion related to probes and diagnostics, and the discussion on antibodies contains discussion related to therapeutic compositions, etc.

B. DEFINITIONS

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[33] The following paragraphs provide a non-exhaustive list of definitions of some of the terms and phrases as used herein. All terms used herein, including those specifically described below in this section, are used in accordance with their ordinary meanings unless the context or definition indicates otherwise. Also unless indicated otherwise, except within

the claims, the use of "or" includes "and" and vice-versa. Non-limiting terms are not to be construed as limiting unless expressly stated (for example, "including" means "including without limitation" unless expressly stated otherwise).

The terms set forth in this application are not to be interpreted in the claims as [34] indicating a "means plus function" relationship unless the word "means" is specifically recited in a claim, and are to be interpreted in the claims as indicating a "means plus function" relationship where the word "means" is specifically recited in a claim. Similarly, the terms set forth in this application are not to be interpreted in method or process claims as indicating a "step plus function" relationship unless the word "step" is specifically recited in the claims, and are to be interpreted in the claims as indicating a "step plus function" relationship where the word "step" is specifically recited in a claim.

"Agonist" indicates a substance, such as a molecule or compound, that interacts with a particular GPCR, for example by binding to the GPCR, to activate, increase, or prolong the amount or the duration of the effect of the biological activity or functionality of 15 the GPCR. Agonists include proteins, nucleic acids, carbohydrates, or any other molecules that bind to and positively modulate the effect of the GPCR. Agonists and other modulators of the particular GPCR can be identified using in vitro or in vivo assays for G protein-coupled receptor expression or G protein-mediated signaling. For example, assays for agonists and other modulators include expressing a particular GPCR in cells or cell membranes, applying putative modulator compounds in the presence or absence of a specific known or putative ligand and then determining the functional effects on the particular GPCR-mediated signaling. Samples or assays comprising a particular GPCR that are treated with a potential agonist or other modulator are compared to control samples without the agonist or other modulator to examine the extent of modulation. Control samples can be assigned a relative 25 activity value for the particular GPCR of 100%. Agonist activity on a particular GPCR is achieved when the G protein-coupled receptor activity value relative to the control is at least about 110%, optionally about 150%, preferably about 200-500%, or about 1000-3000% or higher. Down-modulation (for example by an antagonist) of a particular GPCR is achieved when the particular GPCR activity value relative to the control is at most about 90%, typically about 80%, optionally about 50% or about 25-0% of the 100% value.

[36] "Aggregate," see Complex.

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"Algorithm" refers to a detailed sequence of actions to perform to accomplish some task. In computer programming, refers to instructions given to the computer.

- [38] "Allele" or "allelic sequence" indicates an alternative form of the gene encoding the GPCR. Alleles may result from at least one mutation in the nucleic acid sequence and may result in altered mRNAs or in polypeptides whose structure or function may or may not be altered. Any given natural or recombinant gene may have none, one, or many allelic forms. Common mutational changes that give rise to alleles are generally ascribed to natural deletions, additions, or substitutions of nucleotides. Each of these types of changes may occur alone or in combination with the others, one or more times in a given sequence.
- [39] "Altered" nucleic acid sequences encoding the GPCR include those sequences with 10 deletions, insertions, or substitutions of different nucleotides, resulting in a polynucleotide encoding the same GPCR or a polypeptide variant with at least one substantial structural or functional characteristic of the GPCR. Included within this definition are polymorphisms that may or may not be readily detectable using a particular oligonucleotide probe against the polynucleotide encoding the GPCR. "Altered" proteins may contain deletions, insertions, or substitutions of amino acid residues that produce a silent change and result in a functionally equivalent GPCR. Deliberate amino acid substitutions may be made on the basis of similarity in polarity, charge, solubility, hydrophobicity, hydrophilicity, or the amphipathic nature of the residues, as long as the biological or immunological activity of the GPCR is retained. For example, negatively charged amino acids may include aspartic acid and glutamic acid, positively charged amino acids may include lysine and arginine, and amino acids with uncharged polar head groups having similar hydrophilicity values may include leucine, isoleucine, and valine; glycine and alanine; asparagine and glutamine; serine and threonine; and phenylalanine and tyrosine.
- 25 [40] "Alternative splicing" refers to different ways of cutting and assembling exons to produce mature mRNAs.
 - [41] "Amino acid" refers generally to any of a class of organic compounds that contains at least one amino group, -NH₂, and one carboxyl group, -COOH. The alpha-amino acids, RCH(NH₂)COOH, are the building blocks from which proteins are typically constructed. Amino acid can also refer to artificial chemical analogues or mimetics of a given amino acid as described, depending on the context.

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[42] "Amino acid sequence" refers to a string of amino acids, such as an oligopeptide, peptide, polypeptide, or protein sequence, or a fragment of any of these, including naturally occurring or synthetic molecules and those comprising an artificial chemical analogue or mimetic of a given amino acid. In this context, "biologically active fragments," "biologically functional fragments," "immunogenic fragments," and "antigenic fragments" refer to fragments of the GPCR that are preferably about 15, 25, or 50 or more amino acids in length and that retain a substantial amount of such activity of the GPCR. Where "amino acid sequence" refers to an amino acid sequence of a naturally occurring protein molecule, "amino acid sequence" and like terms are not necessarily limited to the complete native amino acid sequence associated with the recited protein molecule.

- [43] "Amplification" indicates the production of additional copies of something, such as a nucleic acid sequence. Amplification can be generally carried out using polymerase chain reaction (PCR) technologies or other technologies such as the cycling probe reaction (CPR) that are well known in the art. See, e.g., Dieffenbach, C. W. and G. S. Dveksler, PCR Primer, a Laboratory Manual, pp.1-5, Cold Spring Harbor Press, Plainview, N.Y. (1995); U.S. Patents Nos. 5,660,988, 5,731,146 and 6,136,533.
 - [44] "Amplification primers" are oligonucleotides such as natural, analog or artificially created nucleotides that can serve as the basis for the amplification of a selected nucleic acid sequence. They include, for example, both PCR primers and ligase chain reaction oligonucleotides.
- [45] "Analog" or "variant" indicates a GPCR or antigenic peptide that has been modified by deletion, addition, modification, or substitution of one or more amino acid residues compared to the wild-type sequence. Analogs encompass allelic and polymorphic variants, and also muteins and fusion proteins that comprise all or a significant part of such GPCR, e.g., covalently linked via side-chain group or terminal residue to a different protein, polypeptide, or moiety (fusion partner). Variants of a particular GPCR protein refer to an amino acid sequence that is altered by one or more amino acids, for example by one or more amino acid substitution, insertion, deletion or modification, or proteins with or without associated native-pattern glycosylation. The variant may have "conservative" changes. Such "conservative" changes generally are well known in the art and readily determinable for a particular GPCR in view of the present application. Conservative changes include, for example, substitutions where a substituted amino acid has similar structural or chemical

properties to the amino acid it replaced (e.g., negatively charged amino acids include aspartic acid and glutamic acid; positively charged amino acids include lysine, arginine, histidine, asparagine, and glutamine; amino acids containing sulfur include methionine and cysteine; polar hydroxy amino acids include serine, threonine, and tyrosine; large hydrophobic amino acids include phenylalanine and tryptophan; small hydrophobic amino acids include alanine, leucine, isoleucine, and valine). A variant may also have "nonconservative" changes which means that the replacement amino acid provides some substantial change in the amino sequence.

A variant preferably retains at least about 90% identity, and more preferably at least [46] about 95% identity. Within certain embodiments, such variants contain alterations such that the ability of the variant to induce an immunogenic response is not substantially eliminated; in some embodiments the ability to an immunogenic response is not substantially diminished. Modifications of amino acid residues may include but are not limited to aliphatic esters or amides of the carboxyl terminus or of residues containing carboxyl side chains, O-acyl derivatives of hydroxyl group-containing residues, and N-acyl derivatives of the aminoterminal amino acid or amino-group containing residues, e.g., lysine or arginine. Guidance in determining which and how many amino acid residues may be substituted, inserted, deleted or modified without diminishing immunological or biological activity may be found in view of the present application using any of a variety of methods and computer programs known in the art, for example, DNASTAR software. Properties of a variant may generally be evaluated by assaying the reactivity of the variant with, for example, antibodies as described herein or evaluating a biological activity characteristic of the native protein as described herein or as known in the art in view of the present application... Certain polynucleotide variants are capable of hybridizing under appropriately stringent conditions to a naturally occurring DNA sequence encoding a particular GPCR protein (or a complementary sequence). Such hybridizing nucleic acid sequences are also within the scope of this invention.

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[47] "Antagonist" refers to a molecule which interacts with a particular GPCR, for example by binding to the particular GPCR, and prevents, inactivates, decreases or shortens the amount or the duration of the effect of the biological activity of the GPCR. Antagonists include proteins, nucleic acids, carbohydrates, antibodies, or any other molecules that so affect the GPCR. Antagonists can be identified, for example, using appropriate screens

corresponding to those described for agonists above and elsewhere herein or as would be apparent to those skilled in the art in view of the present application.

"Antibody" indicates one type of binding partner, typically encoded by an immunoglobulin gene or immunoglobulin genes, and refers to, for example, intact monoclonal antibodies (including agonist and antagonist antibodies), polyclonal antibodies, phage display antibodies, and multispecific antibodies (e.g., bispecific antibodies) formed, for example, from at least two intact antibodies. Antibody also refers to fragments thereof. which comprise a portion of an intact antibody, generally the antigen-binding or variable region of the intact antibody that are capable of binding the epitopic determinant. Examples 10 of antibody fragments include Fab, Fab', F(ab')2, and Fv fragments, diabodies, linear antibodies, single-chain antibody molecules, and multispecific antibodies formed from antibody fragments. See US Patent No. 6,214,984. Antibody fragments may be synthesized by digestion of an intact antibody or synthesized de novo either chemically or utilizing recombinant DNA technology. Antibodies according to the present invention have at least one of adequate specificity, affinity and capacity to perform the activities desired for the antibodies. Antibodies can, for example, be monoclonal, polyclonal, or combinatorial. Antibodies that bind GPCR polypeptides can be prepared using intact polypeptides or using fragments containing small peptides of interest as the immunizing antigen. The polypeptide or oligopeptide used to immunize an animal (e.g., a mouse, a rat, or a rabbit) can be derived 20 from the translation of RNA, or synthesized chemically, and can be conjugated to a carrier protein if desired. Commonly used carriers that are chemically coupled to peptides include bovine serum albumin, thyroglobulin, and keyhole limpet hemocyanin (KLH). The coupled peptide is then used to immunize the animal.

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- "Antigenic determinant" refers to the antigen recognition site on an antigen (i.e., 25 epitope). Such antigenic determinant may also be immunogenic.
 - "Antisense" refers to any composition containing a nucleic acid sequence that is complementary to a specific nucleic acid sequence. "Antisense strand" refers to a nucleic acid strand that is complementary to the "sense" strand. Antisense molecules may be produced by any method including transcription or synthesis including synthesis by ligating the gene(s) of interest in a reverse orientation to a desired promoter that permits the synthesis of a complementary strand. Once introduced into a cell, the complementary nucleotides can combine with natural sequences produced by the cell to form duplexes and to block either

transcription or translation. The designation "negative" can refer to the antisense strand, and the designation "positive" can refer to the sense strand.

- [51] "Biologically active" or "biologically functional," when referring to an antigenic peptide, indicates that the antigenic peptide induces an immunogenic response specific for the antigenic peptide and thus for the GPCR from which is was obtained. A variant, fragment, etc., of an antigenic peptide is "biologically active" or "biologically functional" if the ability to induce the specific immunogenic response is not substantially diminished. The term "not substantially diminished" means retaining a functionality that is at least about 90% of the functionality of the native antigenic peptide. Appropriate assays designed to evaluate such functionality may be designed based on existing assays known in the art in view of the present application, or on the representative assays provided herein.
- [52] "Annotation" refers to the provision of helpful or identifying information about a GPCR or other open reading frame (ORF), such as locus name, key words, and Medline references.
- 15 [53] "BLAST" refers to the Basic Local Alignment Search Tool, which is a technique for detecting ungapped sub-sequences that match a given query sequence. BLAST can be used as a preliminary step for detecting ORF boundaries.
 - [54] "BLASTP" refers to a BLAST program that compares an amino acid query sequence against a protein sequence database.
- 20 [55] "BLASTX" refers to a BLAST program that compares the six-frame conceptual translation products of a nucleotide query sequence (both strands) against a protein sequence database. BLASTX can be used to create a sub-database of ORFs which may exist on a contig, and to identify the best match between one of these ORFs and a sequence in an external database.
- 25 [56] "Buffer" refers to a component in a solution to provide a buffered solution that resists changes in pH by the action of its acid-base conjugate components.
 - [57] "CDS" refers to the GenBank DNA sequence entry for coding sequence. A coding sequence is a sub-sequence of a DNA sequence that is surmised to encode a gene. A complete gene coding sequence begins with an "ATG" and ends with a stop codon.
- 30 [58] "Clone" in molecular biology refers to a vector carrying an insert DNA sequence.
 - [59] "Cloning" in molecular biology refers to a recombinant DNA technique used to produce multiple, up to millions or more, copies of a DNA sequence. The DNA sequence is

inserted into a small carrier or vector (e.g., plasmid, bacteriophage, or virus) and inserted into a host cell for amplification or expression.

- [60] "Cluster" refers to a group of ORFs related to one another by sequence homology. Clusters are generally determined by a specified degree of homology and overlap (e.g., a stringency).
- [61] "Comparison window" indicates a segment of any one of the number of contiguous positions selected from the group consisting of from 20 to 600, usually about 50 to about 200, more usually about 100 to about 150 in which a sequence may be compared to a reference sequence of the same number of contiguous positions after the two sequences are aligned to enhance sequence similarity. Methods of alignment of sequences for comparison will be readily apparent to a person of ordinary skill in the art in view of the present application.
- [62] "Complementary" or "complementarity" refers to the natural binding of polynucleotides by base pairing. For example, the sequence "A-G-T" binds to the complementary sequence "T-C-A." Complementarity between two single-stranded molecules may be "partial," such that only some of the nucleic acids bind, or it may be "complete," such that all of the nucleotides of at least one of the single-stranded molecules binds to corresponding nucleotides of the other single-stranded molecule. The degree of complementarity between nucleic acid strands has significant effects on the efficiency and strength of the hybridization between the nucleic acid strands. This can be of particular importance in amplification reactions, which can depend upon binding between nucleic acids strands, and in the design and use of peptide nucleic acid (PNA) molecules.
 - [63] "Complex," or "aggregate," indicates a dimer or multimer formed between at least two proteins or other macromolecules, for example a GPCR and its ligand.
 - [64] "Composition" indicates a combination of multiple substances into a mixture.
- 25 [65] "Composition comprising a given amino acid sequence" refers broadly to any composition containing the given amino acid sequence. The composition may comprise a dry formulation, an aqueous solution, or a sterile composition.
 - "Consensus sequence" refers to the sequence that reflects the most common choice of base or amino acid at each position from a series of related DNA, RNA, or protein sequences. Areas of particularly good agreement often represent conserved functional domains. The generation of consensus sequences has typically been subjected to intensive mathematical analysis.

- [67] "Conservative changes" to an amino acid sequence, see Analog.
- [68] "Deletion" refers to a change in the amino acid or nucleotide sequence that results in the absence of one or more amino acid residues or nucleotides.
- [69] "Derivative" refers to chemical modification of an antigenic peptide, or of an antibody specific for and created from the antigenic peptide. A derivative peptide can be modified, for example, by glycosylation or pegylation.
- [70] "Diabodies" refers to one type of antibody comprising small antibody fragments with two antigen-binding sites, which fragments comprise a heavy-chain variable domain (V_H) connected to a light-chain variable domain (V_L) on the same polypeptide chain (V_H-V_L).
- By using a linker that is too short to allow pairing between the two domains on the same chain, the domains pair with the complementary domains of another chain and create two antigen-binding sites. Diabodies are described, for example, in EP 404,097; WO 93/11161; and Holliger et al., Proc. Natl. Acad. Sci. USA, 90:6444-6448 (1993).
- [71] "Database" refers to a structured format for organizing and maintaining information or data, a collection of data records, in a computer-readable form that can be rapidly and easily retrieved. A database is typically stored in a computer-readable memory. Records may comprise web pages, graphics, audio files, text files, or links. Records may or may not be further broken into fields. Database records are usually indexed and come with a search interface to find records of interest.
- 20 [72] "E-value" refers to a result of a FASTA analysis. The number indicates the probability that a match between two sequences is due to random chance.
 - [73] "Expression vector" is a specialized vector constructed so that the gene inserted in the vector can be expressed in the cytoplasm of a host cell.
- [74] "FASTA" refers to a modular set of sequence comparison programs used to compare an amino acid or DNA sequence against all entries in a sequence database. FASTA was written by Professor William Pearson of the University of Virginia Department of Biochemistry. The program uses the rapid sequence algorithm described by Lipman and Pearson (1988) and the Smith-Waterman sequence alignment protocol. FASTA performs a protein to protein comparison.
- 30 [75] "FASTX" refers to a module of the FASTA protocol used to define optimal ORF boundaries while searching for genes. FASTX uses a nucleotide to protein sequence comparison.

- [76] "Fragment," see Portion.
- [77] "GenBank" refers to a family of public databases comprising nucleic acid and amino acid sequence information, including the GenPept bacterial peptide database.
- [78] "Gene" refers to the basic unit of heredity that carries the genetic information for a given RNA or protein molecule. A gene is composed of a contiguous stretch of DNA and contains a coding region that is flanked on each end by regions that are transcribed but not translated. A gene is a segment of DNA involved in producing a biologically active or biologically functional polypeptide chain.
- [79] "Heterologous" indicates a nucleic acid that comprises two or more subsequences that are not found in the same relationship to each other in nature. For instance, the nucleic acid is typically recombinantly produced, having two or more sequences from unrelated genes arranged to make a new functional nucleic acid, e.g., a promoter from one source and a coding region from another source. Similarly, a heterologous protein indicates that the protein comprises two or more subsequences that are not found in the same relationship to each other in nature (e.g., a fusion protein).
 - [80] "Hit Threshold" refers to a pre-set E-value or P-value for evaluating sequence matches. For example, this value can be set at le-6 for finding genes; and at le-15 for clustering genes.
 - [81] "Homology" refers to a degree of complementarity. There may be partial homology or complete homology. The word "identity" may substitute for the word "homology." A partially complementary sequence that at least partially, and substantially, inhibits a corresponding sequence from hybridizing to a target nucleic acid is referred to as "substantially homologous." The inhibition of hybridization of the completely complementary sequence to the target sequence may be examined using a hybridization assay (e.g., Southern or Northern blot, in situ hybridization, solution hybridization) under conditions of reduced stringency. A substantially homologous sequence or hybridization probe will compete for and inhibit the binding of a completely homologous sequence to the target sequence under stringency conditions that inhibit non-specific binding but permit specific binding. The absence of non-specific binding may be tested by the use of a second target sequence which lacks even a partial degree of complementarity (e.g., less than about 30% homology or identity). In the absence of non-specific binding, the substantially

homologous sequence or probe will not hybridize to the second, non-complementary target sequence.

- "Humanized antibody" refers to antibody molecules in which the amino acid [82] sequence in the non-antigen-binding regions has been altered so that the antibody more 5 closely resembles a human antibody, and still retains its original binding ability. Typically, humanized antibodies are human immunoglobulins (recipient antibody) in which residues from a complementarity-determining region (CDR) of the recipient are replaced by residues from a CDR of a non-human species (donor antibody) such as mouse, rat or rabbit having the desired specificity, affinity, and capacity. In some instances, Fv framework residues of the human immunoglobulin are replaced by corresponding non-human residues. Furthermore, humanized antibodies may comprise residues that are found neither in the recipient antibody nor in the imported CDR or framework sequences. These modifications are typically made to further refine and optimize antibody performance. In general, the humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the framework (FR) regions are those of a human immunoglobulin sequence. The humanized antibody optimally also will comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin. For further details see, e.g., Jones et al., Nature, 321:522-525 (1986); 20 Reichmann et al., Nature, 332:323-329 (1988); and, Presta, Curr. Op. Struct. Biol., 2:593-596 (1992).
 - [83] "Identity," see Homology.
 - [84] "Immunocytochemistry" refers to the use of immunologic methods, including a specific antibody, to study cell constituents.
- 25 [85] "Immunohistochemistry" refers to the use of immunologic methods, including a specific antibody, to study specific antigens in tissue slices.
 - [86] "Immunolocalization" refers to the use of immunologic methods, including a specific antibody, to locate molecules or structures within cells or tissues.
- [87] "Immunologically active" refers to the capability of a natural, recombinant, or synthetic GPCR, or any immunogenic fragment thereof, to induce a specific immune response in appropriate animals or cells and to bind with specific antibodies. A polypeptide is "immunologically active" if it is recognized by (e.g., specifically bound by) a B-cell or T-

cell surface antigen receptor. Immunological activity may generally be assessed using well known techniques, such as those summarized in Paul, Fundamental Immunology, 3rd ed., 243-247, Raven Press (1993) and references cited therein. Such techniques include screening polypeptides derived from the native polypeptide for the ability to react with antigen-specific antisera or T-cell lines or clones, which may be prepared in view of the present application using well known techniques. Preferably, an immunologically active portion of a GPCR protein reacts with such antisera or T-cells at a level that is not substantially lower than the reactivity of the full-length polypeptide (e.g., in an ELISA or T-cell reactivity assay). Such screens may generally be performed using methods well known to those of ordinary skill in the art in view of the present application, such as those described in Harlow and Lane, Antibodies: A Laboratory Manual, Cold Spring Harbor Press (1988). B-cell and T-cell epitopes may also be predicted via computer analysis.

- [88] "Immune response" refers to any of the body's immunologic reactions to an antigen such as antibody formation, cellular immunity, hypersensitivity, or immunological tolerance.
- [89] "Insertion" and "addition" when referring to a change in a nucleotide or amino sequence indicate the addition of one or more nucleotides or amino acid residues, respectively, to the sequence.
- [90] "In situ hybridization" refers to use of a nucleic acid probe, typically a DNA or RNA probe, to detect the presence of a DNA or RNA sequence in target cells such as cloned bacterial cells, cultured eukaryotic cells, or tissue samples. In situ hybridization can also be used for locating genes on chromosomes. The process can be performed by preparing a microscope slide with cells in metaphase of mitosis, then treating slide with a weak base to denature the DNA. Next, pour radioactively labeled probe onto the slide under hybridizing conditions, expose the slide to a photographic emulsion for a suitable period such as a few days or weeks, then develop the emulsion.
 - [91] "Isoform" refers to different forms of a protein that may be produced from different genes or from the same gene by alternative RNA splicing.
- [92] "Isolated" generally means that the material is removed from its original environment (e.g., the natural environment if it is naturally occurring).
 - [93] "Library" refers physically to a pool of nucleic acid fragments that has been propagated in a cloning vector. Library can also refer to an electronic collection of genomic

or proteomic sequence data, including raw sequences, contigs, ORFs and loci from a specific organism.

"Ligand" refers to an ion or molecule that binds with another molecule, such as a [94] GPCR, to form a macromolecule such as a receptor-ligand complex. An "endogenous ligand" refers to a native ligand that binds to the receptor of the GPCR and modulates biological activity or functionality of the GPCR in its native environment. A "specific ligand" is a ligand able to bind to a particular GPCR and modulate the biological activity or functionality of the particular GPCR; an endogenous ligand is one example of a specific ligand.

"Microarray" refers to an array of distinct nucleic acid or amino acid molecules [95] arrayed on a substrate, such as paper, nylon or any other type of membrane, filter, chip, glass slide, or any other suitable solid support. Microarrays can also refer to tissue microarrays, composed of small tissue pieces arranged on a slide. U.S. Pat. No. 5,143,854 and PCT Patent Publication Nos. WO 90/15070 and 92/10092.

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"Mimetic" refers to a molecule, e.g., a peptide or non-peptide agent, such as a small molecule, that is able to perform the same biological activity as a certain biologically active agent. For example, some mimetics are molecules comprising the same biological function or activity as the particular GPCR. The structure of the mimetic can be developed from knowledge of the structure of the particular GPCR or portions thereof. For appropriate mimetics, the mimetic is able to effect some or all of the actions of a given antigenic peptide 20 or antibodies against the angtigenic peptide. Such mimetics can be made, in view of the present application, using techniques well known in the art, see, e.g., U.S. Patent Nos. 6,197,752; 6,093,697; 6,207,643; 5,849,323, and can be included in the various processes, methods, and systems, etc., described herein, such as databases, binding partner assays, probes, medicaments, and therapeutics.

"Modulate" refers to controllably changing the activity of a substance or other item, such as the biological activity of a GPCR, antigenic peptide or corresponding antibody. For example, modulation may cause an increase or a decrease in protein activity, binding characteristics, or other biological, functional, or immunological properties of the GPCR.

"Monoclonal antibody" refers to an antibody obtained from a population of 30 [98] substantially homogeneous antibodies, e.g., the individual antibodies comprising the population are identical except for possible naturally occurring mutations that may be present

in minor amounts. Monoclonal antibodies include "chimeric" antibodies (immunoglobulins) in which a portion of the heavy or light chain is identical with or homologous to corresponding sequences in antibodies derived from a particular species or belonging to a particular antibody class or subclass, while the remainder of the chain(s) is identical with or homologous to corresponding sequences in antibodies derived from another species or belonging to another antibody class or subclass, as well as fragments of such antibodies, so long as they exhibit the desired biological activity. U.S. Pat. No. 4,816,567; Morrison et al., P.N.A.S. USA, 81:6851-6855 (1984). Monoclonal antibodies are highly specific, being directed against a single antigenic site. As a matter of distinction, polyclonal antibody 10. preparations typically include different antibodies directed against different determinants (epitopes) of a target antigen whereas each monoclonal antibody is directed against a single determinant on the antigen. Monoclonal antibodies can be synthesized by hybridoma culture. uncontaminated by other immunoglobulins. For example, the monoclonal antibodies to be used in accordance with the present invention may be made by the hybridoma method first 15 described by Kohler and Milstein, Nature, 256:495 (1975), or may be made by recombinant DNA methods. See, e.g., U.S. Pat. No. 4,816,567. Monoclonal antibodies may also be isolated from phage antibody libraries using the techniques described in Clackson et al., Nature, 352:624-628 (1991), and Marks et al., J. Mol. Biol., 222:581-597 (1991), for example. The modifier "monoclonal" indicates the character of the antibody as being 20 obtained from a substantially homogeneous population of antibodies, and is not to be construed as requiring production of the antibody by any particular method.

- [99]. "Nonconservative" changes to an amino acid sequence, see Analog.
- [100]: "Northern blotting" or "Northern analysis" refers to a method used to detect specific RNA sequences. For example, the process can be performed by electrophoresing RNA in a denaturing agarose gel, transferring the gel onto a membrane, and hybridizing with a labeled RNA or DNA probe.
 - [101] "Nucleic acid sequence" refers to a polymer comprising a string of "nucleic acids" such as an oligonucleotide, or a polynucleotide or fragment thereof. The nucleic acid sequence can be from DNA or RNA of genomic or synthetic origin, may be single-stranded or double-stranded, and may represent the sense or the antisense strand. A nucleic acid sequence can also be a PNA or a DNA-like or RNA-like material. Unless stated otherwise,

the term encompasses nucleic acids containing known analogues or mimetics of natural nucleotides that have similar binding properties as the reference nucleic acid.

[102] "Oligonucleotide" refers to a nucleic acid sequence, generally between 6 nucleotides to 60 nucleotides, preferably about 15 to 30 nucleotides, and most preferably about 20 to 25 nucleotides, that can, for example, be used in PCR or other nucleic acid amplification or in a hybridization assay or microarray. "Oligonucleotide" includes "amplimers," "primers," "oligomers," and "probes," as these terms are commonly defined in the art. Oligonucleotides can be chemically synthesized. Such synthetic oligonucleotides may have no 5' phosphate and if so will not ligate to another oligonucleotide without adding a phosphate, typically by using an ATP in the presence of a kinase. A synthetic oligonucleotide will ligate to a fragment that has not been dephosphorylated.

[103] "Operably linked" or "operably connected" indicates that one element of an apparatus, system, or method, etc., is connected to another element of the apparatus, system, or method, etc., such that the two elements are able to perform their intended purposes. For example, when a promoter is linked to a polynucleotide to allow transcription of the polynucleotide, it is "operably linked" to the polynucleotide.

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[104] "Orphan receptor" refers to a receptor for which the endogenous ligand or other ligands inducing biological activity are not known.

[105] "PCR" or "polymerase chain reaction" refers to an *in vitro* method that uses oligonucleotide primers, enzymes, and a series of repetitive temperature cycles to generate millions of copies of a nucleic acid, typically DNA, from an original specimen of a specific DNA sequence, which specimen may be present only in a trace amount.

[106] "Plasmids" refers to extrachromasomal genetic elements composed of DNA or RNA found in both eukaryotic and prokaryotic cells that can propagate themselves autonomously in cells. Plasmids can be used as carriers or vectors to clone DNA molecules. They are designated by a lower case p preceded or followed by capital letters or numbers. The starting plasmids herein are either commercially available, publicly available on an unrestricted basis, or can be constructed from available plasmids in accord with published procedures. In addition, equivalent plasmids to those described are known in the art and will be apparent to the ordinarily skilled artisan in view of the present application.

[107] "Polynucleotide encoding a polypeptide" indicates a polynucleotide that includes only the coding sequence for the polypeptide as well as polynucleotides that include additional coding or non-coding sequence.

- [108] "Portion" or "fragment" with regard to a protein (as in "a portion of a given protein") refers to parts of that protein, a subsequence of the complete amino acid sequence of the receptor containing at least about 8, usually at least about 12, more typically at least about 20, and commonly at least about 30 or more contiguous amino acid residues, up to the entire amino acid sequence minus one amino acid. Thus, a protein "comprising at least a portion of the amino acid sequence of SEQ ID NO:XX" or a protein "comprising at least a portion of the amino acid sequence of a particular GPCR" encompasses the full-length protein and fragments thereof. A portion or fragment of a nucleic acid refers to nucleic acid sequences that are greater than about 12 nucleotides in length, and typically at least about 60 or 100 nucleotides, generally at least about 1000 nucleotides, or at least about 10,000 nucleotides in length, up to the entire nucleic acid sequence minus one nucleic acid.
- 15 [109] "P-value" is a statistical term used to indicate the probability that an event is due to random chance. When used in reference to a result of BLAST searches, the number indicates the probability that a match between two sequences is due to random chance.
- [110] "Receptor" refers to a molecular structure, typically within a cell or on a cell surface, that selectively binds a specific substance (a ligand) and a specific physiologic effect that accompanies the binding. GPCRs are a type of cell-surface receptor, which means a protein in, on, or traversing the cell membrane (in the case of GPCRs, traversing the cell membrane) that recognizes and binds to specific molecules in the surrounding fluid. The binding to a receptor may serve to transport molecules into the cell's interior or to signal the cell to respond in some way.
- 25 [111] "Recombinant" refers to both a method of production and a structure. Some recombinant nucleic acids and proteins are made by the use of recombinant DNA techniques that involve human intervention, either in manipulation or selection. Others are made by fusing two fragments that are not naturally contiguous to each other. Engineered vectors are encompassed, as well as nucleic acids comprising sequences derived using any synthetic oligonucleotide process.
 - [112] "Sample" is used in its usual broad sense. For example, a biological sample suspected of containing nucleic acids encoding the GPCR, or fragments thereof, or the GPCR

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itself, may comprise a bodily fluid; an extract from a cell, chromosome, organelle, or membrane from a cell; a cell; genomic DNA, RNA, or cDNA (in solution or bound to a solid support); a tissue; a tissue print, and the like. Biological sample refers to samples from a healthy individual as well as to samples from a subject suspected of having or susceptible to having, e.g., immune-related diseases, cell growth-related diseases, cell regeneration-related diseases, immunological-related cell proliferative diseases, and autoimmune diseases. Examples of specific diseases include AIDS, allergies, Alzheimer's disease, amyotrophic lateral sclerosis, atherosclerosis, bacterial, fungal, protozoan and viral infections, benign prostatic hypertrophy, bone diseases (e.g., osteoarthritis, osteoporosis), carcinoma (e.g., basal cell carcinoma, breast carcinoma, embryonal carcinoma, ovarian carcinoma, renal cell carcinoma, lung adenocarcinoma, lung small cell carcinoma, pancreatic carcinoma, prostate carcinoma, transitional carcinoma of the bladder, squamous cell carcinoma, thyroid carcinoma), cardiomyopathy, chronic and acute inflammation, circadian rhythm disorders, COPD. Crohn's disease, diabetes, Duchenne muscular dystrophy, embryonal carcinoma, endotoxic shock, environmental stress (e.g., by heat, UV or chemicals), gastrointestinal disorders, glioblastoma multiform, graft vs. host disease, Hodgkin's disease, inflammatory bowel disease, ischemia, stroke, lymphoma, macular degeneration, malignant cytokine production, malignant fibrous histiocytoma, melanoma, meningioma, mesothelioma, multiple sclerosis, nasal congestion, pain, Parkinson's disease, prostate carcinoma, psoriasis, rhabdomyosarcoma, psychotic or neurological disorders (e.g., anxiety, depression, schizophrenia, dementia, mental retardation, memory loss, epilepsy, locomotor problems, respiratory disorders, asthma, eating/body weight disorders including obesity, bulimia, diabetes, anorexia, nausea, hypertension, hypotension), renal disorders, reperfusion injury, rheumatoid arthritis, sarcoma (e.g., chondrosarcoma, Ewing's sarcoma, osteosarcoma), septicemia seminoma sexual/reproductive disorders, tonsil, transitional carcinoma of the bladder, transplant rejection, trauma, tuberculosis, ulcers, ulcerative colitis, urinary retention, vascular and cardiovascular disorders, or any other disease or disorder in which G proteincoupled receptors are involved, as well as learning and/or memory disorders, diabetes, pain perception disorders, anorexia, obesity, hormonal release problems, or any other disease or 30 disorder in which a specific GPCR is involved.

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"Second messengers" refer to intracellular signaling molecules such as cyclic AMP (cAMP), inositol triphosphate, diacylglycerol, or Ca2+. Second messengers, in turn, alter the

activity of other intracellular proteins such as cAMP-dependent protein kinase and Ca²⁺/calmodulin-dependent protein kinases, leading to the transduction and amplification of the original extracellular signal.

[114] "Southern blotting" refers to a method for detecting specific DNA sequences via hybridization. For example, a DNA sample can be electrophoresed in a denaturing agarose gel, transferred onto a membrane, and hybridized with a complementary nucleic acid probe. "Southern" when used in reference to a database indicates an electronic analog of the laboratory technique, which analysis can be used to identify libraries in which a given DNA sequence, such as a gene, EST, or ORF is present. The terms "Northern" and "Western" likewise can be used for electronic analogs to the respective laboratory techniques described above.

[115] "Specific binding" or "specifically binding" refers to an interaction between protein or peptide and a certain substance, such as its specific ligand or antibody, and in some cases its agonists or antagonists. The interaction is dependent upon the presence of a particular structure of the protein recognized by the binding molecule (e.g., the antigenic determinant or epitope). For example, if an antibody specifically binds epitope "A," the presence of a polypeptide containing epitope A or the presence of free unlabeled epitope A will reduce the amount of labeled epitope A that binds to the antibody in a reaction containing free labeled epitope A and the antibody. Conversely, the presence of a polypeptide that does not contain epitope A will not reduce the amount of labeled epitope A that binds to the antibody. Highly specific binding indicates that the protein or peptide binds to its particular ligand, antibody, etc., and does not bind in a significant amount to other proteins present in the sample. Typically, a specific or selective reaction will be at least twice the background signal or noise and more typically more than 10 to 100 times the background signal or noise.

[116] "Stringent conditions" refer to conditions that permit hybridization between complementary polynucleotide sequences. Suitably stringent conditions can be defined by, for example, the concentrations of salt or formamide in the prehybridization and hybridization solutions, or by the hybridization temperature. Stringency can be increased by reducing the concentration of salt, increasing the concentration of formamide, or raising the hybridization temperature. Stringent conditions are dependent upon the type of probe as well as the length of the probe and the GC content of the probe. "Stringent conditions" typically

occur within a range from about Tm-5°C (5°C below the melting temperature (Tm) of the probe) to about Tm-20-25°C for a cRNA probe and to about Tm-15°C for an oligonucleotide "Highly stringent conditions" refers to conditions under which a probe will hybridize to its target sequence, typically in a complex mixture of nucleic acid sequences, but will not substantially hybridize to other sequences. One example of high stringency conditions for a cRNA probe that is 1,000 nucleotides in length and has a GC content of about 60% is about 55-65°C in 50% formamide, 0.1 X SSC, and 200 µg/ml sheared and denatured salmon sperm DNA. One example of low stringency conditions for the same probe in 50% formamide, 0.1 X SSC, and 200 µg/ml sheared and denatured salmon sperm DNA would be 30-35°C. "Very highly stringent conditions" indicates that there must be complete identity between the sequences. The temperature range corresponding to a particular level of stringency can be narrowed further by calculating the purine to pyrimidine ratio of the nucleic acid of interest and adjusting the temperature accordingly. Variations on and modifications of the above ranges and conditions will be readily appreciated by those of skill in the art in view of the present application. As will be understood by those of skill in the art in view of the present application, the stringency of hybridization can be altered to identify or detect identical or related polynucleotide sequences. One guide for nucleic acid hybridization is Tijssen, Laboratory Techniques in Biochemistry and Molecular Biology-v.24 Hybridization with Nucleic Acid Probes, Part I "Overview of principles of hybridization and the strategy of nucleic acid assays" (New York: Elsevier 1993).

[117] "Substantially purified" refers to nucleic acid or amino acid sequences that are removed from their natural environment and are separated from other components from such natural environment, and are at least about 60% free, preferably about 75% or 85% free, and most preferably about 90%, 95% or 99% free from such other components with which they are naturally associated. Substantially purified preferably indicates a substantially homogeneous state and can be in either a dry or aqueous solution or other composition as desired. Purity and homogeneity can be assayed by standard methods, for example on a mass or molar basis, using analytical chemistry techniques such as polyacrylamide gel electrophoresis or high performance liquid chromatography.

[118] "Substitution" when referring to a change in a nucleotide or amino sequence indicates the replacement of one or more nucleotides or amino acids by different nucleotides or amino acids, respectively.

[119] "Variant," see Analog.

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- [120] "Western blotting" or "Western analysis" refers to a method for detecting specific protein sequences. For example, the process can be performed by electrophoresing a protein mixture in a denaturing agarose or acrylamide gel, transferring the mixture onto a membrane, and incubating it with an antibody raised against the protein of interest.
 - [121] Other terms and phrases are defined in other portions of this application.

C. SELECTION OF DESIRED ANTIGENIC PEPTIDES FOR GPCRs AND OTHER POLYPEPTIDES

[122] The present invention provides improved antigenic peptides, for example as set forth in Figure 2, SEQ ID NOS. 692-2292, and improved methods of identifying such antigenic peptides from known or publicly available sequences of polypeptides or proteins, i.e., from a candidate polypeptide sequence. Polypeptide and protein are used in their traditional sense to indicate lengthy amino acid molecules, whereas the antigenic peptide has a length significantly less than the length of the corresponding polypeptide or protein such that the antigenic peptide is capable of providing significantly improved antigenicity relative to the corresponding polypeptide or protein, typically improved specificity, affinity or avidity. The candidate polypeptide can be, for example, a human protein or polypeptide, a naturally occurring protein or polypeptide or a synthetic or recombinant protein or polypeptide.

[123] The antigenic peptides are typically 5 to about 100 amino acids in length, preferably 6 to about 50 amino acids, and further preferably 7 to about 20 amino acids. The antigenic peptides include short antigenic amino acid sequences (i.e., peptides comprising only a portion of an antigenic sequence as set forth in Figure 2 or as identified using the methods described herein, plus an insignificant number of additional amino acids at one or both ends, where insignificant indicates that the extra amino acids do not substantially interfere with the antigenicity of the antigenic peptide). Such short antigenic peptides can be identical to at least 5, 6, 7 or more consecutive amino acids of the sequences herein or identified using the methods described herein, or can have one or two (or more, with increasing length)

conservative amino acid substitution for antigenic peptides comprising more than 6 or 7 consecutive amino acids of the sequences herein or identified using the methods described herein. Antigenic peptides and sequences, and related antibodies and assays and the like, are discussed further elsewhere herein with regard to GPCRs, but such discussions applies to all antigenic peptides produced according to the methods herein, including proteins and polypeptides such as kinases, phosphatases and any other desired protein or polypeptide.

- [124] The identification or selection methods comprise searching the candidate polypeptide sequence using a comparison window of the desired length, then selecting against or rejecting amino acid sequences of the length and having at least 1 characteristic selected from the group consisting of 1) at least two consecutive prolines, 2) at least two consecutive serines, 3) at least two consecutive lysines, 4) at least two consecutive arginines, 5) at least two consecutive aspartic acids, 6) at least two consecutive glutamic acids, 7) methionine, 8) tryptophan, and 9) at least five consecutive amino acids comprising no charged amino acids. Preferably, at least 5, 7, 8, or all of the characteristics are selected.
- 15 [125] The identification or selection methods can also comprise selecting against amino acid sequences having at least 5 consecutive amino acids that are identical to an alternative amino acid sequence from an alternative polypeptide, i.e., some polypeptide other than the candidate polypeptide from which the selected antigen was derived, that is different from the candidate polypeptide, posttranslational modification sites, or highly hydrophobic sequences, which indicates sequences adequately hydrophobic to be located in a lipid membrane such as a cellular membrane. The posttranslational modification sites can be phosphorylation or glycosylation sites.
 - [126] The methods can further comprise performing a BLAST-type or a FAST-type analyses for the candidate polypeptide sequence. Exemplary BLAST-type and FAST-type analyses are described above, including BLAST, BLASTP, BLASTX, FASTA, and FASTX.

D. GENERAL DISCUSSION OF ANTIGENIC PEPTIDES RELATED TO PARTICULAR GPCRS

[127] ANTIGENIC PEPTIDES GENERALLY:

30 [128] The present invention includes antigenic peptides able to induce specific immunogenic responses, and corresponding binding partners. Such antigenic peptides and

binding partners can be cloned, expressed, isolated, purified, and otherwise obtained or manipulated according to routine methods known in the art in view of the present application. The present invention further relates to antigenic peptides having an amino acid [129] sequence from a particular GPCR, including analogs, mimetics, fragments, derivatives, and the like of such antigenic peptides. See SEQ ID NOS. 1-2292, Figures 1-3. The antigenic peptides may be recombinant, natural or synthetic. The antigenic peptides include (i) antigenic peptides in which one or more of the amino acid residues are substituted with a conserved or non-conserved amino acid residue (preferably a conserved amino acid residue) and such substituted amino acid residue may or may not be one encoded by the genetic code, (ii) antigenic peptides in which one or more of the amino acid residues includes a substituent group, (iii) antigenic peptides in which the mature polypeptide is complexed (e.g., fused or otherwise bonded) with another compound, such as a compound to increase the half-life of the polypeptide (for example, polyethylene glycol), and (iv) antigenic peptides in which additional amino acids are fused to the antigenic peptide. Preparing and using such analogs, etc., are within the scope of those skilled in the art in view of the present application. The antigenic peptides additionally include antigenic peptides that have at least about 90% identity to the given antigenic peptide, and preferably at least about 95% identity to the antigenic peptide. The antigenic peptides additionally include antigenic peptides that contain at least five, six, seven or more consecutive amino acids that are identical to the given antigenic peptide, as well as antigenic peptides that contain at least six, seven, eight or more consecutive amino acids that are identical to the given antigenic except for one or two conservative changes within this such stretch of amino acids. The antigenic peptides of the present invention can be produced by peptide synthesis.

[130] EXPRESSION PROFILES BASED ON PROTEINS:

25 [131] An expression profile of a particular GPCR in one or more tissues can be made using antibodies or other binding partners produced using the antigenic peptides herein, then using traditional approaches such as Western blotting, immunohistochemistry analysis, protein array, ligand-binding studies, radioimmunoassay (RIA), and high performance liquid chromatography (HPLC), and immunohistochemistry analysis. H&E staining and other analyses can be used in combination with such immunologically-based analyses.

[132] SCREENING FOR ACTIVITY:

[133] The activity or functionality of an antigenic peptide can be measured using any of a variety of assays known in the art. Similarly, the specificity or affinity of an antibody or other binding partner made using the antigenic peptide can be measured using any of a variety of assays known in the art

The activity or functionality of a particular GPCR may be measured using any of a variety of functional assays in which activation of the receptor in question results in an observable change in the level of some second messenger system, including but not limited to adenylyl cyclase, calcium mobilization, arachidonic acid release, ion channel activity, inositol phospholipid hydrolysis, or guanylyl cyclase. Heterologous expression systems utilizing appropriate host cells to express the nucleic acid of the subject invention are used to obtain the desired second messenger coupling. Receptor activity may also be assayed in an oocyte expression system.

[135] PROTEIN PURIFICATION:

[136] The antigenic peptides and proteins or polypeptides containing them can be purified by standard methods, including but not limited to salt or alcohol precipitation, preparative disc-gel electrophoresis, isoelectric focusing, high pressure liquid chromatography (HPLC), reversed-phase HPLC, gel filtration, cation and anion exchange, partition chromatography, and countercurrent distribution. Suitable purification methods will be readily apparent to those skilled in the art in view of the present application and are disclosed, e.g., in Guide to Protein Purification, Methods in Enzymology, Vol. 182, M. Deutscher, Ed., Academic Press, New York, NY (1990). Purification steps can be followed as part of carrying out assays for ligand binding activity. Particularly where a particular GPCR is being isolated from a cellular or tissue source, it is preferable to include one or more inhibitors of proteolytic enzymes in the assay system, such as phenylmethylsulfonyl fluoride (PMSF).

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- E. CERTAIN ASSAYS, ANTIBODIES, PROBES, THERAPEUTICS, AND OTHER SYSTEMS AND ASPECTS, OF THE INVENTION
 - 1. SYSTEMS AND METHODS FOR SCREENING FOR A PARTICULAR GPCR OR ANTIGENIC PEPTIDE

30 [137] SCREENING FOR ANTIGENIC PEPTIDES:

[138] As noted elsewhere herein, the present invention provides antigenic peptides and antibodies that are specific for a particular GPCR. The invention also provides systems and

methods for using or detecting such peptides, and antibodies against such peptides or corresponding GPCRs in a sample. The assays are based on the detection of the antigenic peptides, typically as they are displayed by the particular GPCR, or the detection of antibodies produced against the particular antigenic peptides and corresponding GPCRs.

5 [139] SCREENING FOR/WITH ANTIGENIC PEPTIDES:

[140] Many assays are characterized by the ability of antigenic peptides for a particular GPCR to be bound by antibodies against them, and the ability of antibodies produced against such antigenic peptides to bind to antigens or epitopes of the particular GPCR in a sample. Some exemplary assays are described below and elsewhere herein.

10 [141] LIST OF ASSAYS:

[142] A variety of assays can detect antibodies that bind specifically to the desired protein in or from a sample, or detect a desired protein bound to one or more antibodies in or from the sample. Exemplary assays are described in detail in Antibodies: A Laboratory Manual, Harlow and Lane (eds.), Cold Spring Harbor Laboratory Press (1988). Representative examples of such assays include: countercurrent immuno-electrophoresis (CIEP), radioimmunoassays, radioimmunoprecipitations, enzyme-linked immunosorbent assays (ELISA), dot blot assays, inhibition or competition assays, sandwich assays, immunostick (dip-stick) assays, simultaneous assays, immunochromatographic assays, immunofiltration assays, latex bead agglutination assays, immunofluorescent assays, biosensor assays, and low-light detection assays. See U.S. Pat. Nos. 4,376,110 and 4,486,530; WO 94/25597; WO/25598.

[143] ENZYME-LINKED IMMUNOSORBENT ASSAYS (ELISA):

[144] One assay for the detection of a particular GPCR is a sandwich assay such as an enzyme-linked immunosorbent assay (ELISA). In one preferred embodiment, the ELISA comprises the following steps: (1) coating the particular GPCR antigenic peptide onto a solid phase, (2) incubating a sample suspected of containing anti-particular GPCR antibodies with the antigenic peptide coated onto the solid phase under conditions that allow the formation of an antigen-antibody complex, (3) adding an anti-antibody (such as anti-IgG) conjugated with a label to be captured by the resulting antigen-antibody complex bound to the solid phase, and (4) measuring the captured label and determining therefrom whether the sample contains anti-particular GPCR antibodies.

[145] IMMUNOFLUORESCENCE ASSAY:

[146] A fluorescent antibody test (FA-test) uses a fluorescently labeled antibody able to bind to one of the proteins of the invention. For detection, visual determinations are made by a technician using fluorescence microscopy, yielding a qualitative result. In one embodiment, this assay is used for the examination of tissue samples or histological sections.

[147] BEAD AGGLUTINATION ASSAYS:

[148] In latex bead agglutination assays, antibodies to one or more of the antigenic peptides of the present invention are conjugated to latex beads. The antibodies conjugated to the latex beads are then contacted with a sample under conditions permitting the antibodies to bind to desired proteins in the sample, if any. The results are then read visually, yielding a qualitative result. In some embodiments, as with certain other assays, this format can be used in the field for on-site testing.

[149] ENZYME IMMUNOASSAYS:

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[150] Enzyme immunoassays (EIA) include a number of different assays that can use the antibodies described in the present application. For example, a heterogeneous indirect EIA uses a solid phase coupled with an antibody of the invention and an affinity purified, anti-IgG immunoglobulin preparation. The solid phase can be a polystyrene microtiter plate. The antibodies and immunoglobulin preparation are then contacted with the sample under conditions permitting antibody binding, which conditions are well known in the art. The results of such an assay can be read visually or using a device such as a spectrophotometer, such as an ELISA plate reader, to yield a quantitative result. An alternative solid phase EIA format includes plastic-coated ferrous metal beads able to be moved during the procedures of the assay by means of a magnet. Yet another alternative is a low-light detection immunoassay format. In this highly sensitive format, the light emission produced by appropriately labeled bound antibodies are quantified automatically. Preferably, the reaction is performed using microtiter plates.

[151] In an alternative embodiment, a radioactive tracer is substituted for the enzyme-mediated detection in an EIA to produce a radioimmunoassay (RIA).

[152] SANDWICH ASSAY:

[153] In a capture-antibody sandwich enzyme assay, the desired protein is bound between an antibody attached to a solid phase, preferably a polystyrene microtiter plate, and a labeled antibody. The results can be measured, for example, using a spectrophotometer, such as an ELISA plate reader.

[154] SEQUENTIAL AND SIMULTANEOUS ASSAYS:

[155] In a sequential assay format, reagents are allowed to incubate with the capture antibody in a stepwise fashion. The test sample is first incubated with the capture antibody. Following a wash step, incubation with the labeled antibody occurs. In a simultaneous assay, the two incubation periods described in the sequential assay are combined. This eliminates one incubation period plus a wash step.

[156] IMMUNOSTICK (DIP-STICK) ASSAYS:

[157] A dipstick/immunostick format is essentially an immunoassay using a polystyrene paddle or dipstick instead of a polystyrene microtiter plate as the solid phase. Reagents are the same and the format can either be simultaneous or sequential.

[158] IMMUNOCHROMATOGRAPHIC ASSAYS:

[159] In a chromatographic strip test format, a capture antibody and a labeled antibody are dried onto a chromatographic strip, which typically comprises nitrocellulose or high porosity nylon bonded to cellulose acetate. The capture antibody is usually spray dried as a line at one end of the strip. At this end, there is an absorbent material that is in contact with the strip. At the other end of the strip, the labeled antibody is deposited in a manner that prevents it from being absorbed onto the membrane. Usually, the label attached to the antibody is a latex bead or colloidal gold. The assay may be initiated by applying the sample immediately in front of the labeled antibody.

20 [160] IMMUNOFILTRATION ASSAYS:

[161] Immunofiltration/immunoconcentration formats combine a large solid-phase surface with directional flow of sample/reagents, which concentrates and accelerates the binding of antigen to antibody. In an exemplary format, the test sample is preincubated with a labeled antibody, and then applied to a solid-phase such as fiber filters, nitrocellulose membranes, or the like. The solid phase can also be precoated with latex or glass beads coated with capture antibody. Detection of analyte is the same as that in a standard immunoassay. The flow of sample/reagents can be modulated by either vacuum or the wicking action of an underlying absorbent material.

[162] BIOSENSOR ASSAYS:

30 [163] A threshold biosensor assay is a sensitive, instrumented assay amenable to screening large numbers of samples at low cost. In one embodiment, such an assay comprises the use of light-addressable potentiometric sensors wherein the reaction involves

the detection of a pH change due to binding of the desired protein by capture antibodies, bridging antibodies, and urease-conjugated antibodies. Upon binding, a pH change is effected that is measurable by translation into electrical potential (µvolts). The assay typically occurs in a very small reaction volume, and is very sensitive; the reported detection limit of the assay is 1,000 molecules of urease per minute.

2. ANTIBODIES

[164] ANTIBODIES GENERATED AGAINST A PARTICULAR ANTIGENIC PEPTIDE AND ITS CORRESPONDING GPCR:

[165] Highly specific, high affinity or antibodies against a particular GPCR or other polypeptide can be generated using the antigenic peptides herein and using antibody generation techniques as described herein or elsewhere. The antibodies produced using the antigenic peptides of the present invention, for example, have a specificity for the corresponding GPCR such that the antibodies can selectively detect the corresponding GPCR in a sample containing non-desired or contaminating proteins or polypeptides, such as a tissue or blood sample. Preferably, the antibodies have a high specificity such that no significant amounts of such proteins or polypeptides are detected, and further preferably have a specificity such that only insubstantial to essentially zero amounts of non-desirable proteins are detected. The antibodies produced using the antigenic peptides of the present invention, for example, typically have an affinity or avidity constant (Ka) of at least about 10⁷ liters/mole, typically a high affinity or avidity at least about 10⁹ liters/mole, preferably at least about 10¹⁰ liters/mole, and further preferably at least about 10¹¹ liters/mole.

[166] The antibodies can be used to conduct immunohistochemistry and other analyses of a variety of tissue samples to determine expression of a particular GPCR in such tissues, for diagnostic assays, and for other desired purposes. The specification will now discuss a variety of antibody types, methods, uses, etc.

[167] ANTIBODIES GENERALLY:

[168] In some embodiments, the present invention provides antibodies and other binding partners created using the antigenic peptides herein and directed to a particular GPCR from which the antigenic peptides were derived. Compositions and uses for such antibodies are contemplated, including diagnostic, medicament, and therapeutic uses. Various diagnostic, medicament, and therapeutic uses for antibodies have been reviewed above and, for example,

in Goldenberg et al., Semin. Cancer Biol., 1(3):217-225 (1990); Beck et al., Semin. Cancer Biol., 1(3):181-188 (1990); Niman, Immunol. Ser., 53:189-204 (1990); Endo, Nippon Igaku Hoshasen Gakkai Zasshi (Japan), 50(8):901-909 (1990); and, U.S. Pat. No. 6,214,984.

[169] Recognized immunoglobulin genes include the kappa, lambda, alpha, gamma, delta, epsilon, and mu constant region genes, as well as myriad immunoglobulin variable region genes. Light chains are classified as either kappa or lambda. Heavy chains are classified as gamma, mu, alpha, delta, or epsilon, which in turn define the immunoglobulin classes, IgG, IgM, IgA, IgD, and IgE, respectively. An exemplary immunoglobulin (antibody) structural unit comprises a tetramer. Each tetramer is composed of two identical pairs of antigenic peptide chains, each pair having one "light" chain (about 25 kD) and one "heavy" chain (about 50-70 kD). The N-terminus of each chain defines a variable region of about 100 to 110 or more amino acids primarily responsible for antigen recognition. The terms variable light chain (V_L) and variable heavy chain (V_H) refer to these light and heavy chains respectively.

15 [170] ANTI-IDIOTYPIC ANTIBODIES:

[171] The present invention encompasses anti-idiotypic antibodies, including polyclonal and monoclonal anti-idiotypic antibodies, that are produced using the antibodies described herein as antigens. These anti-idiotypic antibodies are useful because they may mimic the structures of the antigenic peptides set forth herein.

20 [172] Techniques for producing antibodies, including antibody fragments, include the following.

a. Antibody Preparation

(i) Polyclonal Antibodies

25 [173] ANTIBODY PREP - POLYCLONAL:

[174] Polyclonal antibodies are generally raised in animals by multiple subcutaneous (sc) or intraperitoneal (ip) injections of the relevant antigen and an adjuvant. It may be useful to conjugate the relevant antigen to a protein that is immunogenic in the species to be immunized, e.g., keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, or soybean trypsin inhibitor, using a bifunctional or derivatizing agent, for example, maleimidobenzoyl sulfosuccinimide ester (conjugation through cysteine residues), N-

hydroxysuccinimide (through lysine residues), glutaraldehyde, succinic anhydride, SOCl₂, or R¹N=C=NR, where R and R¹ are different alkyl groups.

[175] ANTIBODY PREP – ADJUVANTS (ALL ABS):

· [176] Suitable adjuvants for the vaccination of animals for the production of polyclonal, monoclonal, and other antibodies include but are not limited to Adjuvant 65 (containing peanut oil, mannide monooleate, and aluminum monostearate); Freund's complete or incomplete adjuvant; mineral gels such as aluminum hydroxide, aluminum phosphate, and hexadecylamine, octadecylamine. surfactants such as lysolecithin. dimethyldioctadecylammonium bromide. N,N-dioctadecyl-N',N'-bis(2-hydroxymethyl) propanediamine, methoxyhexadecylglycerol, and pluronic polyols; polyanions such as pyran, dextran sulfate, poly IC, polyacrylic acid, and carbopol; peptides such as muramyl dipeptide, dimethylglycine, tuftsin, stress proteins, core-containing proteins from a positive stranded RNA virus, see US Pat. No. 6,153,378; and, oil emulsions. The antigenic peptides could also be administered following incorporation into liposomes or other microcarriers.

15 [177] Information concerning adjuvants and various aspects of immunoassays are disclosed, e.g., in the series by P. Tijssen, Practice and Theory of Enzyme Immunoassays, 3rd Edition (1987), Elsevier, New York. Other useful references covering methods for preparing polyclonal antisera include Microbiology, Hoeber Medical Division, Harper and Row (1969); Landsteiner, Specificity of Serological Reactions, Dover Publications, New York (1962); and, Williams, et al., Methods in Immunology and Immunochemistry, Vol. 1, Academic Press, New York (1967).

[178] Animals can be immunized against the antigen, immunogenic conjugates, or derivatives by combining 1 mg or 1 µg of the peptide or conjugate (for rabbits or mice, respectively) with 3 volumes of Freund's complete adjuvant and injecting the solution intradermally at multiple sites. One month later the animals are boosted with 1/5 to 1/10 the original amount of peptide or conjugate in Freund's complete adjuvant by subcutaneous injection at multiple sites. Seven to 14 days later the animals are bled and the serum is assayed for antibody titer. Animals are boosted until the titer plateaus. Preferably, the animal is boosted with the conjugate of the same antigen, but conjugated to a different protein or through a different cross-linking reagent. Conjugates also can be made in recombinant cell culture as protein fusions. In addition, aggregating agents such as alum can be suitably used to enhance the immune response.

(ii) Monoclonal Antibodies

[179] ANTIBODY PREP - MONOCLONAL:

[180] Monoclonal antibodies are obtained from a population of substantially homogeneous antibodies, e.g., the individual antibodies comprising the population are identical except for possible naturally occurring mutations that may be present in minor amounts. For example, monoclonal antibodies can be made using the hybridoma method first described by Kohler and Milstein, Nature, 256:495 (1975), or can be made by recombinant DNA methods, or otherwise as desired.

10 [181] In the hybridoma method, a mouse, or other appropriate host animal, such as a hamster, is immunized as described herein to elicit lymphocytes that produce or are capable of producing antibodies that will bind specifically to the antigenic peptide used for immunization. Alternatively, lymphocytes may be immunized *in vitro*. Lymphocytes then are fused with myeloma cells using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell, Goding, Monoclonal Antibodies: Principles and Practice, pp. 59-103, Academic Press (1986).

[182] The hybridoma cells thus prepared are seeded and grown in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused, parental myeloma cells. For example, if the parental myeloma cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine (HAT medium), which substances prevent the growth of HGPRT-deficient cells.

[183] Preferred myeloma cells are those that fuse efficiently, support stable high-level production of antibody by the selected antibody-producing cells, and are sensitive to a medium such as HAT medium, for example murine myeloma lines, such as those derived from MOPC-21 and MPC-11 mouse tumors available from the Salk Institute Cell Distribution Center, San Diego, CA USA, and SP-2 cells available from the American Type Culture Collection, Rockville, MD USA. Human myeloma and mouse-human heteromyeloma cell lines have also been described for the production of human monoclonal antibodies, Kozbor, J. Immunol., 133:3001 (1984); Brodeur et al., Monoclonal Antibody Production Techniques and Applications, pp. 51-63, Marcel Dekker, Inc., New York (1987).

Culture medium in which hybridoma cells are growing is assayed for production of [184] monoclonal antibodies directed against the antigenic peptide. The binding specificity of antibodies produced by hybridoma cells can be determined by immunoprecipitation or by an in vitro binding assay, such as radioimmunoassay (RIA) or enzyme-linked immunosorbent assay (ELISA). The binding affinity of the monoclonal antibody can, for example, be determined by the Scatchard analysis of Munson and Pollard, Anal. Biochem., 107:220 (1980). The antibodies produced using the antigenic peptides of the present invention, for example, typically have an affinity or avidity constant (Ka) of at least about 10⁷ liters/mole, typically a high affinity or avidity at least about 10⁹ liters/mole. preferably at least about 10¹⁰ liters/mole, and further preferably at least about 10¹¹ liters/mole. [185] After hybridoma cells are identified that produce antibodies of the desired specificity, affinity, or activity, the clones may be subcloned by limiting dilution procedures and grown by standard methods (Goding, supra). Suitable culture media for this purpose include, for example, D-MEM or RPMI-1640 medium. In addition, the hybridoma cells may be grown *in vivo* as ascites tumors in an animal.

The monoclonal antibodies secreted by the subclones are suitably separated from the culture medium, ascites fluid, or serum by conventional immunoglobulin purification A-SEPHAROSETM, for example, protein hydroxyapatite procedures such as, chromatography, gel electrophoresis, dialysis, or affinity chromatography.

20 DNA encoding the monoclonal antibodies can be readily isolated and sequenced using conventional procedures (e.g., by using oligonucleotide probes that are capable of binding specifically to genes encoding the heavy and light chains of murine antibodies). The hybridoma cells serve as a preferred source of such DNA. Once isolated, the DNA may be placed into expression vectors, which can then be transfected into host cells such as E. coli cells, simian COS cells, Chinese hamster ovary (CHO) cells, or myeloma cells that do not otherwise produce immunoglobulin protein, to obtain the synthesis of monoclonal antibodies in the recombinant host cells. Review articles on recombinant expression in bacteria of DNA encoding antibody include Skerra et al., Curr. Opinion in Immunol., 5:256-262 (1993), and Pluckthun, Immunol. Revs., 130:151-188 (1992).

30 [188] **MOABS - COMBINATORIAL:**

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In a further embodiment, antibodies or antibody fragments can be isolated from [189] antibody phage libraries generated using the techniques described in McCafferty et al.,

Nature, 348:552-554 (1990), using the proper antigen such as CD11a, CD18, IgE, or HER-2 to select for a suitable antibody or antibody fragment. Clackson et al., Nature, 352:624-628 (1991) and Marks et al., J. Mol. Biol., 222:581-597 (1991) describe the isolation of murine and human antibodies, respectively, using phage libraries. Subsequent publications describe 5 the production of high affinity (nM range) human antibodies by chain shuffling, Marks et al., Biotechnology, 10:779-783 (1992), as well as combinatorial infection and in vivo recombination as strategies for constructing very large phage libraries. Waterhouse et al., Nuc. Acids. Res., 21:2265-2266 (1993). Combinatorial antibodies are also discussed in Huse et al., Science 246:1275-1281 (1989), and Sastry et al., Proc. Natl. Acad. Sci. USA, 86:5728-5732 (1989), and Alting-Mees et al., Strategies in Molecular Biology 3:1-9 (1990). These references describe a system commercially available from Stratacyte, La Jolla, CA USA. Briefly, mRNA is isolated from a B cell population and utilized to create heavy and light chain immunoglobulin cDNA expression libraries in the \(\lambda \text{IMMUNOZAP(H)} \) and AIMMUNOZAP(L) vectors. These vectors may be screened individually or co-expressed to form Fab fragments or antibodies, see Huse et al., supra, see also Sastry et al., supra. Positive plaques can subsequently be converted to a non-lytic plasmid, which allows for highlevel expression of monoclonal antibody fragments from E. coli.

[190] HUMANIZED MOAB:

[191] Binding partners can also be constructed utilizing recombinant DNA techniques to incorporate the variable regions of a gene that encode a specifically binding antibody. The construction of these binding partners can be readily accomplished by one of ordinary skill in the art in view of the present application. See Larrick et al., Biotechnology, 7:934-938 (1989); Riechmann et al., Nature, 332:323-327 (1988); Roberts et al., Nature, 328:731-734 (1987); Verhoeyen et al., Science 239:1534-1536 (1988); Chaudhary et al., Nature, 339:394-397 (1989); see also U.S. Pat. No. 5,132,405 entitled "Biosynthetic Antibody Binding Sites".) For example, the DNA can be modified by substituting the coding sequence for human heavy- and light-chain constant domains in place of homologous murine sequences, U.S. Pat. No. 4,816,567; Morrison, et al., Proc. Nat. Acad. Sci., 81:6851 (1984), or by covalently joining to the immunoglobulin coding sequence all or part of the coding sequence for a non-immunoglobulin polypeptide. In another example, DNA segments encoding the desired antigen-binding domains specific for the protein or peptide of interest are amplified from appropriate hybridomas and inserted directly into the genome of a cell that produces human

antibodies. See Verhoeyen et al., supra; see also Reichmann et al., supra. Some of these techniques transfer the antigen-binding site of a specifically binding mouse or rat monoclonal antibody or the like to a human antibody. Such antibodies can be preferable for therapeutic use in humans because they are typically not as antigenic as rat or mouse antibodies.

In an alternative embodiment, genes that encode the variable region from a hybridoma producing a monoclonal antibody of interest can be amplified using oligonucleotide primers for the variable region. These primers may be synthesized by one of ordinary skill in the art, or may be purchased from commercially available sources. For instance, primers for mouse and human variable regions including, among others, primers for VHa, VHb, VHC, VHd, CH1, VL and CL regions are available from Stratacyte (La Jolla, CA). These primers may be utilized to amplify heavy- or light-chain variable regions, which may then be inserted into vectors such as IMMUNOZAPTM(H) or IMMUNOZAPTM(L) (Stratacyte), respectively. These vectors may then be introduced into E. coli for expression. Utilizing these techniques, large amounts of a single-chain protein containing a fusion of the VH and VL domains may be produced, see Bird et al., Science 242:423-426 (1988).

[193] ANTIBODY SUBSTITUTIONS - NON-IMMUNOGLOBULIN POLYPEPTIDES (ALL ABS):

[194] Non-immunoglobulin polypeptides can be substituted in monoclonal and other antibodies described herein for the constant domains of an antibody, or they can be substituted for the variable domains of one antigen-combining site of an antibody to create a chimeric bivalent antibody comprising one antigen-combining site having specificity for an antigen and another antigen-combining site having specificity for a different antigen.

[195] CHIMERICS:

[196] Chimeric or hybrid antibodies can also be prepared *in vitro* using known methods in synthetic protein chemistry, including those involving crosslinking agents, in view of the present application. For example, immunotoxins may be constructed using a disulfide-exchange reaction or by forming a thioether bond. Examples of suitable reagents for this purpose include iminothiolate and methyl-4-mercaptobutyrimidate.

[197] ANTIBODY LABELING (ALL ABS):

[198] For diagnostic applications or otherwise as desired, and for monoclonal and other antibodies described herein, the antibodies and other binding partners typically will be labeled with a detectable moiety. The detectable moiety can be any moiety that is capable of

producing, either directly or indirectly, a detectable signal. For example, the detectable moiety may be a radioisotope, such as ³H, ¹⁴C, ³²P, ³⁵S, or ¹²⁵I; a fluorescent or chemiluminescent compound, such as fluorescein isothiocyanate, rhodamine, or luciferin; or an enzyme, such as alkaline phosphatase, beta-galactosidase, or horseradish peroxidase. Any method known in the art for conjugating the antibody or binding partner to the detectable moiety may be employed, including those methods described by Hunter et al., Nature, 144:945 (1962); David et al., Biochemistry, 13:1014 (1974); Pain et al., J. Immunol. Meth., 40:219 (1981); and Nygren, J. Histochem. Cytochem., 30:407 (1982).

(iii) Humanized And Human Antibodies

[199] HUMANIZED AB GENERALLY:

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[200] Methods for humanizing non-human antibodies are well known in the art and have been discussed in part above. Generally, a humanized antibody has one or more amino acid residues introduced into it from a source which is non-human. These non-human amino acid residues are often referred to as "import" residues, which are typically taken from an "import" variable domain. Humanization can be performed essentially following the method of Winter and co-workers, Jones et al., Nature, 321:522-525 (1986); Riechmann et al., Nature, 332:323-327 (1988); Verhoeyen et al., Science, 239:1534-1536 (1988), by substituting rodent CDRs or CDR sequences for the corresponding sequences of a human antibody. Accordingly, such humanized antibodies are chimeric antibodies, U.S. Pat. No. 4,816,567, wherein substantially less than an intact human variable domain has been substituted by the corresponding sequence from a non-human species. In practice, humanized antibodies are typically human antibodies in which some CDR residues and possibly some FR residues are substituted by residues from analogous sites in rodent antibodies.

25 [201] The choice of human variable domains, both light and heavy, to be used in making humanized antibodies is very important to reduce antigenicity. According to the so-called "best-fit" method, the sequence of the variable domain of a rodent antibody is screened against the entire library of known human variable-domain sequences. The human sequence that is closest to that of the rodent is then accepted as the human framework (FR) for the humanized antibody. Sims et al., J. Immunol., 151:2296 (1993); Chothia and Lesk, J. Mol. Biol., 196:901 (1987). Another method uses a particular framework derived from the consensus sequence of all human antibodies of a particular subgroup of light or heavy chains.

The same framework may be used for several different humanized antibodies. Carter et al., Proc. Natl. Acad. Sci. USA, 89:4285 (1992); Presta et al., J. Immunol., 151:2623 (1993).

[202] It is typically desirable that antibodies be humanized with retention of high affinity for the antigen and other favorable biological properties. To achieve this goal, according to one method, humanized antibodies are prepared by a process of analysis of the parental sequences and various conceptual humanized products using three-dimensional models of the parental and humanized sequences. Three-dimensional immunoglobulin models are commonly available and are familiar to those skilled in the art. Computer programs are available that illustrate and display probable three-dimensional conformational structures of selected candidate immunoglobulin sequences. Inspection of these displays permits analysis of the likely role of the residues in the functioning of the candidate immunoglobulin sequence, e.g., the analysis of residues that influence the ability of the candidate immunoglobulin to bind antigen. In this way, FR residues can be selected and combined from the consensus and import sequences so that the desired antibody characteristic, such as increased affinity for the target antigen(s), is achieved. In general, CDR residues are directly and most substantially involved in influencing antigen binding.

[203] It is also possible to produce transgenic animals (e.g., mice) that are capable, upon immunization, of producing a full repertoire of human antibodies in the absence of endogenous immunoglobulin production. For example, it has been described that the homozygous deletion of the antibody heavy-chain joining region (J_H) gene in chimeric and germ-line mutant mice results in complete inhibition of endogenous antibody production. Transfer of the human germ-line immunoglobulin gene array in such germ-line mutant mice will result in the production of human antibodies upon antigen challenge. See, e.g., Jakobovits et al., Proc. Natl. Acad. Sci. USA. 90:2551-255 (1993); Jakobovits et al., Nature, 362:255-258 (1993); Bruggemann et al., Year Immuno., 7:33 (1993). Human antibodies can also be produced in phage-display libraries, Hoogenboom and Winter, J. Mol. Biol., 227:381 (1991); Marks et al., J. Mol. Biol., 222:581 (1991).

(iv) Antibody Fragments

30 [204] ANTIBODY FRAGMENTS:

[205] Various techniques have been developed for the production of antibody fragments. Such fragments can be derived via proteolytic digestion of intact antibodies, see, e.g.,

Morimoto et al., J. Biochem. Biophys. Meth. 24:107-117 (1992) and Brennan et al., Science, 229:81 (1985). Fragments can also be produced directly by recombinant host cells. For example, antibody fragments can be isolated from antibody phage libraries discussed above. Fab'-SH fragments can be directly recovered from *E. coli* and chemically coupled to form F(ab')₂ fragments, Carter et al., Biotechnology 10:163-167 (1992). F(ab')₂ fragments can be isolated directly from recombinant host cell culture. Other techniques for the production of antibody fragments will be apparent to the skilled practitioner.

(v) Bispecific Antibodies

10 [206] BISPECIFIC ANTIBODIES GENERALLY:

[207] Bispecific antibodies (BsAbs) are antibodies that have binding specificities for at least two different antigens. Bispecific antibodies can be derived from full-length antibodies or from antibody fragments, e.g., F(ab')₂ bispecific antibodies.

[208] Methods for making bispecific antibodies are known in the art. Traditional production of full-length bispecific antibodies is based on the coexpression of two immunoglobulin heavy chain-light chain pairs, where the two chains have different specificities, Millstein and Cuello, Nature, 305:537-539 (1983). Because of the random assortment of immunoglobulin heavy and light chains, these hybridomas (quadromas) produce a mixture of potentially 10 different antibody molecules, of which only one has the correct bispecific structure. Purification of the correct molecule, which is usually accomplished by affinity chromatography steps, is rather cumbersome, and the product yields are low. Similar procedures are disclosed in WO 93/08829, and in Traunecker et al., E.M.B.O. J., 10:3655-3659 (1991).

25 binding specificities (antibody-antigen combining sites) are fused to immunoglobulin constant domain sequences. The fusion is preferably with an immunoglobulin heavy chain constant domain, comprising at least part of the hinge, C_H 2, and C_H 3 regions. It is preferred to have the first heavy-chain constant region (C_H 1) containing the site necessary for light chain binding, present in at least one of the fusions. DNAs encoding the immunoglobulin heavy chain fusions and, if desired, the immunoglobulin light chain, are inserted into separate expression vectors, and are co-transfected into a suitable host organism. This provides for great flexibility in adjusting the mutual proportions of the three polypeptide fragments in

embodiments when unequal ratios of the three polypeptide chains used in the construction provide the improved yields. It is, however, possible to insert the coding sequences for two or all three polypeptide chains in one expression vector when the expression of at least two polypeptide chains in equal ratios results in high yields or when the ratios are of no particular significance.

[210] ANTIBODIES - HYBRID IMMUNOGLOBULIN HEAVY CHAIN:

[211] In one embodiment of this approach, the bispecific antibodies are composed of a hybrid immunoglobulin heavy chain with a first binding specificity in one arm, and a hybrid immunoglobulin heavy chain-light chain pair (providing a second binding specificity) in the other arm. This asymmetric structure may facilitate the separation of the desired bispecific compound from unwanted immunoglobulin chain combinations, as the presence of an immunoglobulin light chain in only one half of the bispecific molecule provides for a facile method of separation. This approach is discussed in WO 94/04690. For further details of generating bispecific antibodies see, for example, Suresh et al., Meth. Enzymol., 121:210 (1986).

[212] ANTIBODIES - CROSS-LINKED OR "HETEROCONJUGATE":

[213] Bispecific antibodies include cross-linked or "heteroconjugate" antibodies. For example, one of the antibodies in the heteroconjugate can be coupled to avidin, the other to biotin. Such antibodies have, for example, been proposed to target immune system cells to unwanted cells, U.S. Pat. No. 4,676,980), and for treatment of HIV infection, WO 91/00360, WO 92/200373, and EP 03089). Heteroconjugate antibodies may be made using any convenient cross-linking methods. Suitable cross-linking agents are well known in the art, and are disclosed in U.S. Pat. No. 4,676,980, along with a number of cross-linking techniques.

25 [214] ANTIBODIES - DIABODIES:

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[215] The "diabody" technology described by Hollinger et al., Proc. Natl. Acad. Sci. USA, 90:6444-6448 (1993) has provided an alternative mechanism for making BsAb fragments. The fragments comprise a heavy-chain variable domain (V_H) connected to a light-chain variable domain (V_L) by a linker that is too short to allow pairing between the two domains on the same chain. Accordingly, the V_H and V_L domains of one fragment are forced to pair with the complementary V_L and V_H domains of another fragment, thereby forming two antigen-binding sites.

[216] Another strategy for making BsAb fragments by the use of single-chain Fv (sFv) dimers has also been reported. See Gruber et al., J. Immunol., 152:5368 (1994). These researchers designed an antibody comprising the V_H and V_L domains of a first antibody joined by a 25-amino-acid-residue linker to the V_H and V_L domains of a second antibody. The refolded molecule bound to fluorescein and the T-cell receptor and redirected the lysis of human tumor cells that had fluorescein covalently linked to their surface.

[217] ANTIBODIES - OTHER:

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- [218] Techniques for generating bispecific antibodies from antibody fragments have also been described in the literature. For example, bispecific antibodies can be prepared using chemical linkage. Brennan et al., Science, 229:81 (1985) describe a procedure wherein intact antibodies are proteolytically cleaved to generate F(ab')₂ fragments. These fragments are reduced in the presence of the dithiol complexing agent sodium arsenite to stabilize vicinal dithiols and prevent intermolecular disulfide formation. The Fab' fragments generated are then converted to thionitrobenzoate (TNB) derivatives. One of the Fab'-TNB derivatives is then reconverted to the Fab'-thiol by reduction with mercaptoethylamine and is mixed with an equimolar amount of the other Fab'-TNB derivative to form the BsAb. The BsAbs produced can be used as agents for the selective immobilization of enzymes.
- [219] Fab'-SH fragments can be directly recovered from E. coli, which can be chemically coupled to form bispecific antibodies. Shalaby et al., J. Exp. Med., 175:217-225 (1992) describe the production of a fully humanized BsAb F(ab')₂ molecule. Each Fab' fragment was separately secreted from E. coli and subjected to directed chemical coupling in vitro to form the BsAb. The BsAb thus formed was able to bind to cells overexpressing the HER2 receptor and normal human T cells, as well as trigger the lytic activity of human cytotoxic lymphocytes against human breast tumor targets. See also Rodriguez et al., Int. J. Cancers (Suppl.) 7:45-50 (1992).
- [220] Various techniques for making and isolating BsAb fragments directly from recombinant cell culture have also been described. For example, bispecific F(ab')₂ heterodimers have been produced using leucine zippers. Kostelny et al., J. Immunol., 148(5):1547-1553 (1992). The leucine zipper peptides from the Fos and Jun proteins are linked to the Fab' portions of two different antibodies by gene fusion. The antibody homodimers are reduced at the hinge region to form monomers and then re-oxidized to form the antibody heterodimers.

b. Antibody Purification

[221] ANTIBODY PURIFICATION GENERALLY:

[222] When using recombinant techniques, the antibody can be produced intracellularly, in the periplasmic space, or directly secreted into the medium. If the antibody is produced intracellularly, as a first step, the particulate debris, either host cells or lysed fragments, is removed, for example, by centrifugation or ultrafiltration. Carter et al., Bio/Technology 10:163-167 (1992), describe a procedure for isolating antibodies which are secreted to the periplasmic space of *E. coli*. Briefly, cell paste is thawed in the presence of sodium acetate (pH 3.5), EDTA, and phenylmethylsulfonylfluoride (PMSF) over about 30 min. Cell debris can be removed by centrifugation. Where the antibody is secreted into the medium, supernatants from such expression systems are generally first concentrated using a commercially available protein concentration filter, for example, an Amicon or Millipore Pellicon ultrafiltration unit. A protease inhibitor such as PMSF may be included in any of the foregoing steps to inhibit proteolysis and antibiotics may be included to prevent the growth of adventitious contaminants.

[223] BEFORE LPHIC:

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The antibody composition prepared from the cells is preferably subjected to at least [224] one purification step prior to LPHIC. Examples of suitable purification steps include hydroxyapatite chromatography, gel electrophoresis, dialysis, and affinity chromatography. The suitability of protein A as an affinity ligand depends on the species and isotype of any immunoglobulin Fc domain that is present in the antibody. Protein A can be used to purify antibodies that are based on human $\gamma 1$, $\gamma 2$, or $\gamma 4$ heavy chains, Lindmark et al., J. Immunol. Meth. 62:1-13 (1983). Protein G has been recommended for mouse isotypes and for human γ3, Guss et al., E.M.B.O. J., 5:1567-1575 (1986). The matrix to which the affinity ligand is attached is often agarose, but other matrices are available. Mechanically stable matrices such as controlled pore glass or poly(styrenedivinyl)benzene allow for faster flow rates and shorter processing times than can be achieved with agarose. Where the antibody comprises a C_H 3 domain, the Bakerbond ABXTM resin (J. T. Baker, Phillipsburg, N.J.) is useful for purification. Other techniques for protein purification such as fractionation on an ionexchange column, ethanol precipitation, Reverse Phase HPLC, chromatography on silica. chromatography on heparin SEPHAROSETM, chromatography on an anion or cation

exchange resin (such as a polyaspartic acid column), chromatofocusing, SDS-PAGE, and ammonium sulfate precipitation are also available depending on the antibody to be recovered.

[225] LPHIC:

[226] Following any preliminary purification step(s), the mixture comprising the antibody of interest and contaminant(s) can be subjected to LPHIC. See US Patent No. 6,214,984. Often, the antibody composition to be purified will be present in a buffer from the previous purification step. However, it may be necessary to add a buffer to the antibody composition prior to the LPHIC step. Many buffers are available and can be selected by routine experimentation. The pH of the mixture comprising the antibody to be purified and at least one contaminant in a loading buffer is adjusted to a pH of about 2.5-4.5 using either an acid or base, depending on the starting pH. The loading buffer can have a low salt concentration (e.g., less than about 0.25 M salt).

[227] The mixture is loaded on the HIC column. HIC columns normally comprise a base matrix (e.g., cross-linked agarose or synthetic copolymer material) to which hydrophobic ligands (e.g., alkyl or aryl groups) are coupled. One example of an HIC column comprises an agarose resin substituted with phenyl groups (e.g., a Phenyl SEPHAROSETM column). Many HIC columns are available commercially. Examples include, but are not limited to, Phenyl SEPHAROSE 6 FAST FLOWTM column with low or high substitution (Pharmacia LKB Biotechnology, AB, Sweden); Phenyl SEPHAROSETM High Performance column (Pharmacia LKB Biotechnology, AB, Sweden); Octyl SEPHAROSETM High Performance column (Pharmacia LKB Biotechnology, AB, Sweden); FRACTOGELTM EMD Propyl or FRACTOGELTM EMD Phenyl columns (E. Merck, Germany); MACRO-PREPTM Methyl or MACRO-PREPTM t-Butyl Supports (Bio-Rad, California); WP HI-Propyl (C₃)TM column (J. T. Baker, New Jersey); and TOYOPEARLTM ether, phenyl, or butyl columns (TosoHaas, PA).

[228] The antibody is typically eluted from the column using an elution buffer that is the same as the loading buffer. The elution buffer can be selected using routine experimentation in view of the present application. The pH of the elution buffer may be between about 2.5-4.5 and have a low salt concentration (e.g., less than about 0.25 M salt). It may not be necessary to use a salt gradient to elute the antibody of interest; the desired product may be recovered in the flow-through fraction that does not bind significantly to the column.

[229] The LPHIC step provides a way to remove a correctly folded and disulfide bonded antibody from unwanted contaminants (e.g., incorrectly associated light and heavy fragments). The method can provide an approach to substantially remove an impurity characterized as a correctly folded antibody fragment whose light and heavy chains fail to associate through disulfide bonding. Antibody compositions prepared using LPHIC can be up to about 95% pure or more. Purities of more than about 98% have been reported. US Patent No. 6,214,984.

[230] POST LPHIC:

[231] Antibody compositions prepared by LPHIC can be further purified as desired using techniques which are well known in the art. Diagnostic or therapeutic formulations of the purified protein can be made by providing the antibody composition in a physiologically acceptable carrier, examples of which are provided below. To remove contaminants (e.g., unfolded antibody and incorrectly associated light and heavy fragments) from the HIC column so that it can be re-used, a composition including urea (e.g., 6.0 M urea, 1% MES buffer pH 6.0, 4 mM ammonium sulfate) can be flowed through the column.

c. Some Uses For Antibodies Described Herein

(i) Generally

[232] GENERALLY:

20 [233] The present invention comprises any suitable use for the antibodies and other binding partners discussed herein. The following provides some of the desired uses, including diagnostic and therapeutic uses. Various diagnostic and therapeutic uses for antibodies have been reviewed in Goldenberg et al., Semin. Cancer Biol., 1(3):217-225 (1990); Beck et al., Semin. Cancer Biol., 1(3):181-188 (1990); Niman, Immunol. Ser. 53:189-204 (1990); and, Endo, Nippon Igaku Hoshasen Gakkai Zasshi (Japan) 50(8):901-909 (1990), for example.

[234] ASSAYS:

[235] The antibodies can be used in immunoassays, such as enzyme immunoassays. BsAbs can be useful for this type of assay; one arm of the BsAb can be designed to bind to a specific epitope on the enzyme so that binding does not cause enzyme inhibition, the other arm of the antibody can be designed to bind to an immobilizing matrix ensuring a high enzyme density at the desired site. Examples of such diagnostic BsAbs include those having

specificity for IgG as well as ferritin, and those having binding specificities for horseradish peroxidase (HRP) as well as a hormone, for example. Monoclonal and polyclonal antibodies are also exemplary antibodies for immunoassays.

[236] The antibodies can be designed for use in two-site immunoassays. For example, two antibodies are produced binding to two separate epitopes on the analyte protein; one antibody binds the complex to an insoluble matrix, the other binds an indicator enzyme.

[237] **DIAGNOSTIC USES:**

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Antibodies can also be used for immunodiagnosis, in vitro or in vivo or otherwise, [238] of various diseases or conditions based on the presence or absence of a particular GPCR. Such diseases and conditions include, e.g., immune-related diseases, cell growth-related diseases, cell regeneration-related diseases, immunological-related cell proliferative diseases, and autoimmune diseases. Examples of specific diseases include AIDS, allergies, Alzheimer's disease, amyotrophic lateral sclerosis, atherosclerosis, bacterial, fungal, protozoan and viral infections, benign prostatic hypertrophy, bone diseases (e.g., osteoarthritis, osteoporosis), carcinoma (e.g., basal cell carcinoma, breast carcinoma, embryonal carcinoma, ovarian carcinoma, renal cell carcinoma, lung adenocarcinoma, lung small cell carcinoma, pancreatic carcinoma, prostate carcinoma, transitional carcinoma of the bladder, squamous cell carcinoma, thyroid carcinoma), cardiomyopathy, chronic and acute inflammation, circadian rhythm disorders, COPD, Crohn's disease, diabetes, Duchenne 20 muscular dystrophy, embryonal carcinoma, endotoxic shock, environmental stress (e.g., by heat, UV or chemicals), gastrointestinal disorders, glioblastoma multiform, graft vs. host disease, Hodgkin's disease, inflammatory bowel disease, ischemia, stroke, lymphoma, macular degeneration, malignant cytokine production, malignant fibrous histiocytoma, melanoma, meningioma, mesothelioma, multiple sclerosis, nasal congestion, pain, 25 Parkinson's disease, prostate carcinoma, psoriasis, rhabdomyosarcoma, psychotic or neurological disorders (e.g., anxiety, depression, schizophrenia, dementia, mental retardation, memory loss, epilepsy, locomotor problems, respiratory disorders, asthma, eating/body weight disorders including obesity, bulimia, diabetes, anorexia, nausea, hypertension, hypotension), renal disorders, reperfusion injury, rheumatoid arthritis, sarcoma (e.g., 30 chondrosarcoma, Ewing's sarcoma, osteosarcoma), septicemia. sexual/reproductive disorders, tonsil, transitional carcinoma of the bladder, transplant rejection, trauma, tuberculosis, ulcers, ulcerative colitis, urinary retention, vascular and

cardiovascular disorders, or any other disease or disorder in which G protein-coupled receptors are involved, as well as learning and/or memory disorders, diabetes, pain perception disorders, anorexia, obesity, hormonal release problems, or any other disease or disorder in which a specific GPCR is involved.

[239] To facilitate this diagnostic use, an antibody that binds a particular GPCR, when such is differentially expressed in tumors or other target diseases, can be conjugated with a detectable marker (e.g., a chelator that binds a radionuclide). Examples of tumor-associated antigens being used in a similar fashion include an antibody having specificity for the tumor-associated antigen CEA used for imaging colorectal and thyroid carcinomas and the anti-p185^{HER2} antibody used for detecting cancers characterized by amplification of the HER2 protooncogene. Other uses for the antibodies of the present invention will be apparent to the skilled practitioner in view of the present application.

(ii) Assays

15 [240] ASSAYS:

- [241] For certain applications such as some diagnostic and other assay applications, the antibody typically can be labeled directly or indirectly with a detectable moiety. The detectable moiety can be any moiety that is capable of producing, either directly or indirectly, a detectable signal. For example, the detectable moiety may be a radioisotope, such as ³H, ¹⁴C, ³²P, ³⁵S, or ¹²⁵I; a fluorescent or chemiluminescent compound, such as fluorescein isothiocyanate, rhodamine, or luciferin; or an enzyme, such as alkaline phosphatase, beta-galactosidase, or HRP.
- [242] Any method known in the art for separately conjugating the antibody to the detectable moiety may be employed, including those methods described by Hunter et al., Nature, 144:945 (1962); David et al., Biochemistry, 13:1014 (1974); Pain et al., J. Immunol. Meth. 40:219 (1981); and, Nygren, J. Histochem. and Cytochem. 30:407 (1982).
- [243] The antibodies of the present invention may be employed in any desired assay method, such as competitive binding assays, direct, and indirect sandwich assays, and immunoprecipitation assays. Zola, Monoclonal Antibodies: A Manual of Techniques, pp.
- 30 147-158 (CRC Press, Inc. (1987).

[244] COMPETITIVE BINDING ASSAYS:

[245] Competitive binding assays rely on the ability of a labeled standard to compete with the test sample analyte for binding with a limited amount of antibody. The amount of analyte in the test sample is inversely proportional to the amount of standard that becomes bound to the antibody. To facilitate determining the amount of standard that becomes bound, the antibody generally is insolubilized before or after the competition, so that the standard, and analyte that are bound to the antibody may conveniently be separated from the standard, and analyte which remain unbound.

[246] BsAbs are particularly useful for sandwich assays which involve the use of two molecules, each capable of binding to a different immunogenic portion, or epitope, of the sample to be detected. In a sandwich assay, the test sample analyte is bound by a first arm of the antibody which is immobilized on a solid support, and thereafter a second arm of the antibody binds to the analyte, thus forming an insoluble three part complex. See, e.g., U.S. Pat. No. 4,376,110. The second arm of the antibody may itself be labeled with a detectable moiety (direct sandwich assays) or may be measured using an anti-immunoglobulin antibody that is labeled with a detectable moiety (indirect sandwich assay). For example, one type of sandwich assay is an ELISA assay, in which case the detectable moiety is an enzyme. Assays are discussed further elsewhere herein in relation to binding partners such as antibodies, and antigenic peptides for particular GPCRs, including assays searching for or using such antigenic peptides, and would be apparent to those skilled in the art in view of the present application.

(iii) Affinity Purification

[247] AFFINITY PURIFICATION:

[248] The antibodies also are useful for the affinity purification of an antigen of interest such as a particular GPCR from sources such as recombinant cell culture or natural sources.

(iv) Therapeutics

[249] THERAPEUTIC USES:

[250] Therapeutic compositions, and uses, etc., for the antibodies described herein will now be discussed. As with other parts of this application, this section does not contain the entire discussion of therapeutic uses or compositions, etc., for antibodies, other sections discuss both antibodies, and therapeutics, and the discussion in this section applies to certain

other aspects discussed herein. Turning to antibodies and therapeutics, the antibodies can be used, for example, for redirected cytotoxicity (e.g., to kill tumor cells), as a vaccine adjuvant, for delivering thrombolytic agents to clots, for delivering immunotoxins to tumor cells, for converting enzyme activated prodrugs at a target site (e.g., a tumor), for treating infectious diseases or targeting immune complexes to cell surface receptors.

[251] THERAPEUTIC FORMULATIONS:

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[252] Therapeutic formulations of the antibody can be prepared for storage by mixing the antibody having the desired degree of purity with optional physiologically acceptable carriers, excipients, or stabilizers (Remington's Pharmaceutical Sciences, 16th edition, Osol, A., Ed. (1980), for example in the form of lyophilized cake or aqueous solutions. Acceptable carriers, excipients, or stabilizers are nontoxic to recipients at the dosages, and concentrations employed, and include buffers such as phosphate, citrate, and other organic acids; antioxidants including ascorbic acid; low molecular weight (less than about 10 residues) polypeptides; proteins, such as serum albumin, gelatin, or immunoglobulins; hydrophilic polymers such as polyvinylpyrrolidone; amino acids such as glycine, glutamine, asparagine, arginine, or lysine; monosaccharides, disaccharides, and other carbohydrates including glucose, mannose, or dextrins; chelating agents such as EDTA; sugar alcohols such as mannitol or sorbitol; salt-forming counterions such as sodium; or nonionic surfactants such as Tween, Pluronics, or polyethylene glycol (PEG).

20 [253] The antibodies also may be entrapped in microcapsules prepared, for example, by interfacial polymerization (for coacervation techniques or bv poly-[methylmethacrylate] hydroxymethylcellulose or gelatin-microcapsules, and microcapsules, respectively), in colloidal drug delivery systems (for example, liposomes, albumin microspheres, microemulsions, nano-particles, and nanocapsules), or in macroemulsions. Such techniques are disclosed in Remington's Pharmaceutical Sciences, 25 supra.

[254] THERAPEUTIC FORMULATIONS -STERILE:

[255] An antibody to be used for *in vivo* human administration should be sterile. This can be accomplished by filtration through sterile filtration membranes, for example prior to or following lyophilization and reconstitution. The antibody ordinarily will be stored in lyophilized form or in solution. Therapeutic antibody compositions generally are placed into

a container having a sterile access port, for example, an intravenous solution bag or vial having a stopper pierceable by a hypodermic injection needle.

[256] THERAPEUTIC ADMINISTRATIONS:

poly-D-(-)-3-hydroxybutyric acid, EP 133,988.

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[257] The route of antibody administration is in accord with known methods, e.g., injection or infusion by intravenous, intraperitoneal, intracerebral, intramuscular, intraocular, intraarterial, or intralesional routes, or by sustained release systems as noted below.

[258] The antibody can be administered, for example, continuously by infusion or by bolus injection. Suitable examples of sustained-release preparations include semipermeable matrices of solid hydrophobic polymers containing the protein, which matrices are in the form of shaped articles, e.g., films, or microcapsules. Examples of sustained-release matrices include polyesters, hydrogels (e.g., poly(2-hydroxyethyl-methacrylate) as described by Langer et al., J. Biomed. Mater. Res., 15:167-277 (1981), and Langer, Chem. Tech., 12:98-105 (1982), or poly(vinylalcohol)), polylactides, U.S. Pat. No. 3,773,919, EP 58,481, copolymers of L-glutamic acid and gamma ethyl-L-glutamate, Sidman et al., Biopolymers, 22:547-556 (1983), non-degradable ethylene-vinyl acetate, Langer et al., supra, degradable lactic acid-glycolic acid copolymers such as the LUPRON DEPOTTM (injectable microspheres composed of lactic acid-glycolic acid copolymer and leuprolide acetate), and

[259] THERAPEUTIC ADMINISTRATIONS – SUSTAINED RELEASE-POLYMERS:

[260] While polymers such as ethylene-vinyl acetate and lactic acid-glycolic acid sustain release of molecules for over 100 days, certain hydrogels release proteins for shorter time periods. When encapsulated antibodies remain in the body for a long time, they may denature or aggregate as a result of exposure to moisture at 37°C, resulting in a loss of biological activity and possible changes in immunogenicity. Rational strategies can be devised for antibody stabilization depending on the mechanism involved. For example, if the aggregation mechanism is discovered to be intermolecular S--S bond formation through thiodisulfide interchange, stabilization may be achieved by modifying sulfhydryl residues, lyophilizing from acidic solutions, controlling moisture content, using appropriate additives, and developing specific polymer matrix compositions.

[261] THERAPEUTIC ADMINISTRATIONS – SUSTAINED RELEASE-LIPOSOMES:

[262] Sustained-release antibody compositions also include liposomally entrapped antibody. Liposomes containing the antibody can be prepared by methods such as those in DE 3,218,121; Epstein et al., Proc. Natl. Acad. Sci. USA, 82:3688-3692 (1985); Hwang et al., Proc. Natl. Acad. Sci. USA, 77:4030-4034 (1980); EP 52,322; EP 36,676; EP 88,046; EP 143,949; EP 142,641; Japanese patent application 83-118008; U.S. Pat. Nos. 4,485,045 and 4,544,545; and EP 102,324. Ordinarily the liposomes are of the small (about 200-800 Angstroms) unilamellar type in which the lipid content is greater than about 30 mol. % cholesterol, the selected proportion being adjusted for the optimal antibody therapy.

[263] THERAPEUTICALLY EFFECTIVE AMOUNT:

[264] An effective amount of antibody to be employed therapeutically will depend, for example, upon the therapeutic objectives, the route of administration, and the condition of the patient. Accordingly, it will be necessary for the therapist to titer the dosage and modify the route of administration as required to obtain the optimal therapeutic effect. A typical daily dosage might range from about 1 µg/kg to up to 10 mg/kg or more, depending on the factors mentioned above. Typically, the clinician will administer antibody until a dosage is reached that achieves the desired effect. The progress of this therapy is easily monitored by conventional assays.

5. DRUG DESIGN BASED ON THE ANTIGENS HEREIN OR ANTIBODIES THERETO

[265] DISEASE/CONDITIONS LIST:

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[266] The peptides and antibodies of the present invention can serve as valuable tools for designing drugs for treating various pathophysiological conditions such as immune-related diseases, cell growth-related diseases, cell regeneration-related diseases, immunological-related cell proliferative diseases, and autoimmune diseases. Examples of specific diseases include AIDS, allergies, Alzheimer's disease, amyotrophic lateral sclerosis, atherosclerosis, bacterial, fungal, protozoan and viral infections, benign prostatic hypertrophy, bone diseases (e.g., osteoarthritis, osteoporosis), carcinoma (e.g., basal cell carcinoma, breast carcinoma, embryonal carcinoma, ovarian carcinoma, renal cell carcinoma, lung adenocarcinoma, lung small cell carcinoma, pancreatic carcinoma, prostate carcinoma, transitional carcinoma of the bladder, squamous cell carcinoma, thyroid carcinoma), cardiomyopathy, chronic and acute inflammation, circadian rhythm disorders, COPD, Crohn's disease, diabetes, Duchenne

muscular dystrophy, embryonal carcinoma, endotoxic shock, environmental stress (e.g., by heat, UV or chemicals), gastrointestinal disorders, glioblastoma multiform, graft vs. host disease, Hodgkin's disease, inflammatory bowel disease, ischemia, stroke, lymphoma, macular degeneration, malignant cytokine production, malignant fibrous histiocytoma, melanoma, meningioma, mesothelioma, multiple sclerosis, nasal congestion, pain, Parkinson's disease, prostate carcinoma, psoriasis, rhabdomyosarcoma, psychotic or neurological disorders (e.g., anxiety, depression, schizophrenia, dementia, mental retardation, memory loss, epilepsy, locomotor problems, respiratory disorders, asthma, eating/body weight disorders including obesity, bulimia, diabetes, anorexia, nausea, hypertension, hypotension), renal disorders, reperfusion injury, rheumatoid arthritis, sarcoma (e.g., chondrosarcoma, Ewing's sarcoma, osteosarcoma), septicemia, seminoma, sexual/reproductive disorders, tonsil, transitional carcinoma of the bladder, transplant rejection, trauma, tuberculosis, ulcers, ulcerative colitis, urinary retention, vascular and cardiovascular disorders, or any other disease or disorder in which G protein-coupled receptors are involved, as well as learning and/or memory disorders, diabetes, pain perception disorders, anorexia, obesity, hormonal release problems, or any other disease or disorder in which a specific GPCR is involved or that would be readily apparent to those skilled in the art in view of the present application.

EXAMPLES

20 [267] The Examples below provide information as follows: Example 1 relates to the identification and selection of the antigens set forth in Figure 2. Examples 2 to 4 relate to antibody production and purification based on such antigens. Examples 5 to 10 relate to H&E staining. And, Example 11 relates to Western blot analyses.

EXAMPLE 1: SELECTION OF ANTIGENS

[268] Antigenic peptides were derived from the amino acid sequence of a particular GPCR based on analyses of likely antigen-containing regions and specificity of those regions for the protein/gene of interest. The specificity of the antigen peptides (approximately 20 amino acids in length) for antibody generation was determined using the outlined techniques, including BLAST of several public databases. These public databases included but were not limited to GenBank, Swiss Prot Human, Swiss Prot NonHuman, GenPeptH, GenPept M, and

LifeSpan's proprietary databases. With respect to specificity, parameters that precluded the use of a particular peptide included the presence of 6 or more contiguous amino acids with sequence identity to protein(s) other than the protein of interest, the presence of sites of posttranslational modification, including phosphorylation and glycosylation, and highly hydrophobic sequences, which could indicate potential in situ localization within the plasma membrane. The peptides were analyzed for antigenicity using the published algorithm of Hopp, T. P., and Woods, K. R, Proc. Natl. Acad. Sci. U.S.A. 78, 3824-3828, (1981). Additional considerations in antigenic peptide design included 1) selection against sequences with multiple prolines in a row, 2) selection against sequences with multiple serines in a row, 3) selection against sequences with multiple lysines in a row, 4) selection against sequences with multiple arginines in a row 5) selection against sequences with multiple aspartic acids in a row, 6) selection against sequences with multiple glutamic acids in a row, 7) selection against peptides containing methionine or tryptophan, which can become oxidized as a result of the cyclization reaction, and 8) avoidance of stretches of 5 or more amino acids having no uncharged amino acids (which also resulted in a desirable charge to peptide length ratio of at least 1 charge:5 residues). The selected antigenic peptides are set forth in the Sequence Listing and in Figure 2.

EXAMPLE 2: ANTIBODY PRODUCTION SCHEDULE

- 20 [269] Day 0 Pre-immune serum collection (approximately 5.0 ml). Immunize using 200 µg antigen peptide per rabbit in Complete Freund's Adjuvant.
 - [270] Day 14 Immunize using 100 µg antigen per rabbit in Incomplete Freund's Adjuvant.
- [271] Day 28 Immunize using 100 µg antigen per rabbit in Incomplete Freund's Adjuvant.
 - [272] Day 42 Immunize using 100 µg antigen per rabbit in Incomplete Freund's Adjuvant.
 - [273] Day 49 First production bleed; obtain 24.0 26.0 ml.
- [274] Day 56 Immunize using 100 μg antigen per rabbit in Incomplete Freund's 30 Adjuvant.
 - [275] Day 63 Second production bleed and ELISA analysis.

[276] Day 70 - Immunize using 100 µg antigen per rabbit in Incomplete Freund's Adjuvant.

[277] Day 77 - Third production bleed and affinity purification.

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EXAMPLE 3: IMMUNOSORBENT PURIFICATION OF ANTISERUM: COUPLING OF PEPTIDE TO CNBR-ACTIVATED SEPHAROSE 4B

[278] Weigh out 0.8 g of CNBr-activated Sepharose 4B (2.5 ml of final gel volume). Wash and re-swell on sintered glass filter with 1 mM HCl, followed by coupling buffer (0.1 M NaHCO₃, 0.25 M NaCl, pH 8.5). Dissolve 10 mg of protein or peptide in coupling buffer. Mix protein solution with gel suspension and incubate 2 hours at room temperature or overnight at 4°C. Block remaining active groups with 0.2 M glycine buffer, pH 8.1. Wash away excess adsorbed protein with coupling buffer, followed by 0.1 M acetate buffer containing 0.5 M NaCl, pH 4.3. Equilibrate the column with phosphate-buffered saline (PBS), pH 7.7.

EXAMPLE 4: IMMUNOSORBENT PURIFICATION OF ANTISERUM: AFFINITY PURIFICATION OF ANTISERUM

[279] Dilute 10 ml of clear antiserum 1:1 with PBS, pH 7.7, apply to affinity column at a flow rate of 0.3 ml/minute, and monitor absorbance of eluate at 280 nm. Collect fractions of unbound material and rinse column with PBS, pH 7.7. Elute bound antibody with 0.2 M glycine, pH 1.85, and collect eluate until absorbance at 280 nm returns to baseline. Neutralize all collected fractions with 1 M Tris-HCl, pH 8.5 immediately after collection. Determine OD at 280 nm, and determine the total OD recovered. Conduct ELISA analysis with the corresponding antigen to confirm the presence and identity of recovered antibody and the removal of all antibody from the original serum. Concentrate antibody to approximately 2.0 mg/ml and dialyze against PBS with 0.01% NaN₃.

EXAMPLE 5: PREPARATION OF ANTIBODY DILUTIONS

[280] The purpose of this protocol is to dilute antibodies in solution. Materials include Tris-HCL Buffer with carrier protein and 0.015 M NaN₃ (Dako Antibody Diluent #S0809 (DAKO, Carpentaria, CA); vials containing the antibodies described above or commercial antibodies against the particular GPCR; pipetmen and disposable tips; container of chopped ice; 12 ml Dako reagent tubes; and, reagent tube rack.

[281] The procedure is a) calculate proportions of antibody and diluent according to desired concentrations and volume requirements; b) label reagent tubes and place in rack; c) pipette needed volume of diluent into tube(s); d) place vials of antibodies into ice; e) invert and/or flick antibody vial(s) 3 or 4 times to insure suspension; f) pipette required volume of antibody(s) into corresponding diluent volumes; and, g) mix gently.

EXAMPLE 6: PREPARATION OF AUTOSTAINER SOLUTIONS

[282] The purpose of this protocol is the preparation of concentrated solutions for use in a DAKO autostainer. Materials include DAKO[®] TBST (Tris Buffered Saline Containing Tween-S3306), 10X Concentrate, DAKO[®] Target Retrieval Solution, 10x Concentrate (S1699), deionized H₂O, 20L container, with lid, marked at the 10L level, DAKO[®] TBS (Tris Buffered Saline-S1968), and DAKO Tween[®] (S1966).

[283] The procedure to make TBST 10x Concentrate is a) pour 2 500 ml bottles DAKO[®] TBST into a 20 L container, b) add deionized H₂O until solution level is at 10 L mark, c) replace lid and shake 10 to 20 times, d) pour diluted DAKO[®] TBST into autostainer carboy(s) as designated. The procedure to make Target Retrieval Solution is a) measure 135 ml of deionized H₂O and pour into slide bath, b) measure 15 ml of DAKO[®] Target Retrieval solution, c) add to H₂O, and d) agitate. This solution is then used in the steam method of target retrieval, Example 9, below. The procedure to make TBS is a) fill 20L container to 10L mark with deionized H₂O, b) add 2 envelopes of DAKO[®] TBS, c) add 5 ml of DAKO TWEEN[®], and d) replace lid and agitate 10 to 20 times.

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EXAMPLE 7: PREPARATION OF SOLUTIONS FOR ANTIBODY DETECTION

[284] Solutions for antibody detection are prepared using Vector[®] Biotinylated antibody (BA series), Vectastain[®] ABC-AP Kit (AK-5000), 10 mM sodium phosphate, pH 7.5, 0.9% saline (PBS), Vector[®] Red Alkaline Phosphatase Substrate Kit I (SK-5100), and 100 mM Tris-HCl, pH 8.2 Buffer. To prepare biotinylated antibody, add 10 ml of PBS to reagent tube, add 1 drop biotinylated antibody to the PBS, then mix gently. To prepare ABC, to 10 ml of PBS, add 2 drops each of Reagent A and Reagent B, mix immediately, then allow to stand 30 minutes before use. To prepare AP Red, which should be prepared immediately

before use, to 5 ml of Tris-HCl buffer, add 2 drops of Reagent 1 and mix well, add 2 drops of Reagent 2 and mix well, then add 2 drops of Reagent 3 and mix well.

EXAMPLE 8: DEPARAFFINIZATION AND REHYDRATION OF SAMPLES

[285] The purpose of this protocol is to remove paraffin from and rehydrate preserved tissues in preparation for IHC procedures. Materials and equipment include fume hood, vertical slide rack(s), three xylene (VWR #72060-088) baths, three 100% alcohol blend (VWR #72060-050) baths, two 95% alcohol blend (VWR #72060-052) baths, one 70% alcohol blend (VWR #72060-056) bath, and Tris-Buffered Saline (DAKO* \$1968) + Tween* (DAKO \$1966).

[286] Insert the slides into the vertical rack(s). Move slides through baths inside fume hood as follows:

Xylene 5 Minutes
Xylene 5 Minutes
Xylene 5 Minutes
100% Alcohol 2 Minutes
100% Alcohol 2 Minutes
100% Alcohol 1 Minute

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95% Alcohol 2 Minutes 95% Alcohol 2 Minutes

70% Alcohol 1 Minute

[287] Finally, place slides into a container with TBST.

EXAMPLE 9: STEAM METHOD OF TARGET RETRIEVAL

[288] The purpose of this protocol is to optimize antibody binding within paraffin embedded tissues. Materials and equipment included a steamer, deionized H₂O, target retrieval solution, 10X concentrate (DAKO #S1699), 250 ml graduated cylinder, 15 ml graduated cylinder, staining dish(es), and deparaffinized and rehydrated tissue on microscope slides in immersed TBST. The procedure is to a) fill the steamer with deionized H₂O to appropriate depth as indicated, b) turn the steamer on, c) in a graduated cylinder, measure 135ml of deionized H₂O and pour into staining dish(es), d) pipette 15ml of target retrieval solution and release into deionized H₂O, e) place the staining dish(es) into the basket of the steamer and heat for at least 10 minutes to preheat, f) add rack(s) containing tissue slides to heated target retrieval solution, g) cover and steam for 20 minutes, h) remove container from

steamer and let stand at room temperature for 20 minutes, i) transfer rack(s) with slides to container(s) of TBST, and j) slides are now ready for staining procedures.

EXAMPLE 10: ANTIBODY DETECTION

[289] The deparaffinized, rehydrated, and steamed (if needed) slides are loaded onto racks within a DAKO autostainer and then the autostainer is run according to the manufacturer's instructions. The slides are removed and the autostainer is turned off.

EXAMPLE 11: WESTERN BLOTTING

10 [290] The purpose of this protocol is to visualize the immunoreactivity of the antibodies described above against the particular GPCR on a western blot. Materials and equipment included western blot membrane, TBS Tween (TBST: 100 mM Tris-HCl pH 7.5, 150 mM NaCl, 0.1% TweenTM 20), 5% non-fat dried milk in TBST (blotto), antibody of interest (primary), peroxidase-conjugated AffiniPure goat anti-rabbit IgG (H+L) (secondary) – Jackson ImmunoResearch, ECL solution (Amersham Biosciences, Uppsala Sweden), film, developer D-19, fixer, rocking platform.

[291] During the blotting procedure, the blot is kept wet at all times and on a substantially level surface. The Western blot is placed right-side up in 10 ml of blotto. The membrane is flipped over and the dish rocked so that the solution covered it. The membrane is then flipped back to the right side and solution is again rocked over it. The blot is then placed on a shaker for at least 1 hour. Ten ml of primary antibody are prepared by diluting 1:500 in blotto.

[292] The blotto is removed from the Western blot and replaced with the primary antibody. The blot is flipped again and placed on the shaker for 1 hour. Secondary antibody and peroxidase-conjugated AffiniPure goat anti-rabbit IgG (H+L) are prepared 1:20,000 in 10 ml of blotto. The primary antibody is removed and the Western blot is washed 3 times with 10 ml of blotto. The blotto is removed and replaced with the secondary antibody solution. The blot is flipped and placed on the shaker for 1 hour. The secondary antibody is removed and the blot washed 2 times with 10 ml of blotto. The blotto is removed and the blot is washed 2 times with 10 ml of blotto. The blotto is removed and the blot is a prepared by combining equal amounts of Solution 1 and 2.

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[293] The blotto is removed and 1 ml of ECL is placed on the blot. The blot is flipped and let sit for 1 minute. The blot is placed on plastic wrap and immediately covered with plastic wrap. The ECL is pressed out. The blot is placed on the film, then the film is developed.

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[294] From the foregoing, it will be appreciated that, although specific embodiments of the invention have been described herein for purposes of illustration, various modifications may be made without deviating from the spirit and scope of the invention. Accordingly, the invention includes all permutations and combinations of the subject matter set forth herein and is not limited except as by the appended claims.

WHAT IS CLAIMED IS:

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1. An isolated antigenic peptide according to any one of SEQ ID NOS. 692-2292.

- 2. An isolated antigenic peptide comprising an amino acid sequence that is at least about 90% identical to a sequence set forth in any one of SEQ ID NOS. 692-2292.
- 3. An isolated antigenic peptide that is an analog of an antigenic peptide according to any one of SEQ ID NOS. 692-2292.
- 4. An isolated antigenic peptide comprising a short antigenic amino acid sequence that is identical to at least 5 consecutive amino acids set forth in any one of SEQ ID NOS. 692-2292.
 - 5. An isolated antigenic peptide comprising a short antigenic amino acid sequence that is identical to or contains no more than one conservative amino acid substitution over at least 7 consecutive amino acids set forth in any one of SEQ ID NOS. 692-2292.
 - 6. A kit for the detection of antibodies against a particular GPCR in a sample comprising:
 - a) an isolated antigenic peptide according to any one of claims 1-5 and derived from the particular GPCR, and
- 20 b) at least one of a reagent or a device for detecting the antibodies.
- 7. An isolated antibody having high specificity and high affinity or avidity for a particular GPCR comprising a peptide sequence that is identical to any one of SEQ ID NOS. 692-703, 713-730, 744-802, 807-820, 825-875, 880-889, 917-941, 950-964, 971-984, 989-993, 1010-1013, 1021-1024, 1029-1043, 1049-1052, 1057-1072, 1087-1113, 1124-1151, 1161-1172, 1179-1187, 1198-1209, 1228-1231, 1245-1257, 1271-1279, 1304-1308, 1369-1372, wherein the antibody was produced using an isolated antigenic peptide comprising the peptide sequence that is identical to the any one of SEQ ID NOS. 692-703, 713-730, 744-802, 807-820, 825-875, 880-889, 917-941, 950-964, 971-984, 989-993, 1010-1013, 1021-1024, 1029-1043, 1049-1052, 1057-1072, 1087-1113, 1124-1151, 1161-1172, 1179-1187, 30 1198-1209, 1228-1231, 1245-1257, 1271-1279, 1304-1308, 1369-1372.
 - 8. An isolated antibody having high specificity and high affinity or avidity for a particular GPCR comprising a peptide sequence that is at least about 90% identical to any

one of SEQ ID NOS. 692-703, 713-730, 744-802, 807-820, 825-875, 880-889, 917-941, 950-964, 971-984, 989-993, 1010-1013, 1021-1024, 1029-1043, 1049-1052, 1057-1072, 1087-1113, 1124-1151, 1161-1172, 1179-1187, 1198-1209, 1228-1231, 1245-1257, 1271-1279, 1304-1308, 1369-1372, wherein the antibody was produced using the peptide sequence that is at least about 90% identical to the any one of SEQ ID NOS. 692-703, 713-730, 744-802, 807-820, 825-875, 880-889, 917-941, 950-964, 971-984, 989-993, 1010-1013, 1021-1024, 1029-1043, 1049-1052, 1057-1072, 1087-1113, 1124-1151, 1161-1172, 1179-1187, 1198-1209, 1228-1231, 1245-1257, 1271-1279, 1304-1308, 1369-1372.

- 9. An isolated antibody having high specificity and high affinity or avidity for a particular GPCR comprising a peptide sequence that is an analog to any one of SEQ ID NOS. 692-703, 713-730, 744-802, 807-820, 825-875, 880-889, 917-941, 950-964, 971-984, 989-993, 1010-1013, 1021-1024, 1029-1043, 1049-1052, 1057-1072, 1087-1113, 1124-1151, 1161-1172, 1179-1187, 1198-1209, 1228-1231, 1245-1257, 1271-1279, 1304-1308, 1369-1372, wherein the antibody was produced using an isolated antigenic peptide comprising the peptide sequence that is the analog to the any one of SEQ ID NOS. 692-703, 713-730, 744-802, 807-820, 825-875, 880-889, 917-941, 950-964, 971-984, 989-993, 1010-1013, 1021-1024, 1029-1043, 1049-1052, 1057-1072, 1087-1113, 1124-1151, 1161-1172, 1179-1187, 1198-1209, 1228-1231, 1245-1257, 1271-1279, 1304-1308, 1369-1372.
- 10. An isolated antibody having high specificity and high affinity or avidity for a particular GPCR comprising a peptide sequence that is identical to at least 5 consecutive amino acids set forth any one of SEQ ID NOS. 692-703, 713-730, 744-802, 807-820, 825-875, 880-889, 917-941, 950-964, 971-984, 989-993, 1010-1013, 1021-1024, 1029-1043, 1049-1052, 1057-1072, 1087-1113, 1124-1151, 1161-1172, 1179-1187, 1198-1209, 1228-1231, 1245-1257, 1271-1279, 1304-1308, 1369-1372, wherein the antibody was produced using a short isolated antigenic peptide comprising the at least 5 consecutive amino acids set forth in the any one of SEQ ID NOS. 692-703, 713-730, 744-802, 807-820, 825-875, 880-889, 917-941, 950-964, 971-984, 989-993, 1010-1013, 1021-1024, 1029-1043, 1049-1052, 1057-1072, 1087-1113, 1124-1151, 1161-1172, 1179-1187, 1198-1209, 1228-1231, 1245-1257, 1271-1279, 1304-1308, 1369-1372.
- 11. An isolated antibody specific for a particular GPCR comprising a peptide sequence that is identical to any one of SEQ ID NOS. 704-712, 731-743, 774-777, 803-806, 821-824, 876-879, 890-916, 942-949, 965-970, 985-988, 994-1009, 1014-1020, 1025-1028,

1044-1048, 1053-1056, 1073-1086, 1114-1123, 1152-1160, 1173-1178, 1188-1197, 1210-1227, 1232-1244, 1258-1270, 1280-1303, 1309-1368, 1373-1377, 1386-1389, 1394-1402, 1462-1482, 1496-1525, 1542-1549, 1557-1563, 1583-1649, 1656-1679, 1684-1688, 1693-1732, 1744-1752, 1765-1839, 1846-1854, 1855-1866, 1871-1917, 1926-1941, 1952-1955, 1960-1980, 1985-2141, 2152-2165, and 2170-2292, wherein the antibody was produced using an isolated antigenic peptide comprising the peptide sequence that is identical to the any one of SEQ ID NOS. 704-712, 731-743, 774-777, 803-806, 821-824, 876-879, 890-916, 942-949, 965-970, 985-988, 994-1009, 1014-1020, 1025-1028, 1044-1048, 1053-1056, 1073-1086, 1114-1123, 1152-1160, 1173-1178, 1188-1197, 1210-1227, 1232-1244, 1258-1270, 1280-1303, 1309-1368, 1373-1377, 1386-1389, 1394-1402, 1462-1482, 1496-1525, 1542-1549, 1557-1563, 1583-1649, 1656-1679, 1684-1688, 1693-1732, 1744-1752, 1765-1839, 1846-1854, 1855-1866, 1871-1917, 1926-1941, 1952-1955, 1960-1980, 1985-2141, 2152-2165, and 2170-2292.

- 12. An isolated antibody specific for a particular GPCR comprising a peptide sequence that is at least about 90% identical to any one of SEQ ID NOS. 704-712, 731-743, 774-777, 803-806, 821-824, 876-879, 890-916, 942-949, 965-970, 985-988, 994-1009, 1014-1020, 1025-1028, 1044-1048, 1053-1056, 1073-1086, 1114-1123, 1152-1160, 1173-1178, 1188-1197, 1210-1227, 1232-1244, 1258-1270, 1280-1303, 1309-1368, 1373-1377, 1386-1389, 1394-1402, 1462-1482, 1496-1525, 1542-1549, 1557-1563, 1583-1649, 1656-1679, 1684-1688, 1693-1732, 1744-1752, 1765-1839, 1846-1854, 1855-1866, 1871-1917, 1926-1941, 1952-1955, 1960-1980, 1985-2141, 2152-2165, and 2170-2292, wherein the antibody was produced using the peptide sequence that is at least about 90% identical to the any one of SEO ID NOS. 704-712, 731-743, 774-777, 803-806, 821-824, 876-879, 890-916, 942-949, 965-970, 985-988, 994-1009, 1014-1020, 1025-1028, 1044-1048, 1053-1056, 1073-1086, 1114-1123, 1152-1160, 1173-1178, 1188-1197, 1210-1227, 1232-1244, 1258-1270, 1280-25 1303, 1309-1368, 1373-1377, 1386-1389, 1394-1402, 1462-1482, 1496-1525, 1542-1549, 1557-1563, 1583-1649, 1656-1679, 1684-1688, 1693-1732, 1744-1752, 1765-1839, 1846-1854, 1855-1866, 1871-1917, 1926-1941, 1952-1955, 1960-1980, 1985-2141, 2152-2165, and 2170-2292.
- 30 13. An isolated antibody specific for a particular GPCR comprising a peptide sequence that is an analog to any one of SEQ ID NOS. 704-712, 731-743, 774-777, 803-806, 821-824, 876-879, 890-916, 942-949, 965-970, 985-988, 994-1009, 1014-1020, 1025-1028,

1044-1048, 1053-1056, 1073-1086, 1114-1123, 1152-1160, 1173-1178, 1188-1197, 1210-1227, 1232-1244, 1258-1270, 1280-1303, 1309-1368, 1373-1377, 1386-1389, 1394-1402, 1462-1482, 1496-1525, 1542-1549, 1557-1563, 1583-1649, 1656-1679, 1684-1688, 1693-1732, 1744-1752, 1765-1839, 1846-1854, 1855-1866, 1871-1917, 1926-1941, 1952-1955, 1960-1980, 1985-2141, 2152-2165, and 2170-2292, wherein the antibody was produced using an isolated antigenic peptide comprising the peptide sequence that is the analog to the any one of SEQ ID NOS. 704-712, 731-743, 774-777, 803-806, 821-824, 876-879, 890-916, 942-949, 965-970, 985-988, 994-1009, 1014-1020, 1025-1028, 1044-1048, 1053-1056, 1073-1086, 1114-1123, 1152-1160, 1173-1178, 1188-1197, 1210-1227, 1232-1244, 1258-1270, 1280-1303, 1309-1368, 1373-1377, 1386-1389, 1394-1402, 1462-1482, 1496-1525, 1542-1549, 1557-1563, 1583-1649, 1656-1679, 1684-1688, 1693-1732, 1744-1752, 1765-1839, 1846-1854, 1855-1866, 1871-1917, 1926-1941, 1952-1955, 1960-1980, 1985-2141, 2152-2165, and 2170-2292.

- 14. An isolated antibody specific for a particular GPCR comprising a peptide 15 sequence that is identical to at least 5 consecutive amino acids set forth any one of SEO ID NOS. 704-712, 731-743, 774-777, 803-806, 821-824, 876-879, 890-916, 942-949, 965-970, 985-988, 994-1009, 1014-1020, 1025-1028, 1044-1048, 1053-1056, 1073-1086, 1114-1123, 1152-1160, 1173-1178, 1188-1197, 1210-1227, 1232-1244, 1258-1270, 1280-1303, 1309-1368, 1373-1377, 1386-1389, 1394-1402, 1462-1482, 1496-1525, 1542-1549, 1557-1563, 1583-1649, 1656-1679, 1684-1688, 1693-1732, 1744-1752, 1765-1839, 1846-1854, 1855-1866, 1871-1917, 1926-1941, 1952-1955, 1960-1980, 1985-2141, 2152-2165, and 2170-2292, wherein the antibody was produced using a short isolated antigenic peptide comprising the at least 5 consecutive amino acids set forth in the any one of SEO ID NOS: 704-712, 731-743, 774-777, 803-806, 821-824, 876-879, 890-916, 942-949, 965-970, 985-988, 994-1009, 1014-1020, 1025-1028, 1044-1048, 1053-1056, 1073-1086, 1114-1123, 1152-1160, 1173-1178, 1188-1197, 1210-1227, 1232-1244, 1258-1270, 1280-1303, 1309-1368, 1373-1377, 1386-1389, 1394-1402, 1462-1482, 1496-1525, 1542-1549, 1557-1563, 1583-1649, 1656-1679, 1684-1688, 1693-1732, 1744-1752, 1765-1839, 1846-1854, 1855-1866, 1871-1917, 1926-1941, 1952-1955, 1960-1980, 1985-2141, 2152-2165, and 2170-2292.
- 30 15. A kit for the detection of antibodies against the particular GPCR of claim 5 comprising:
 - a) an isolated antibody according to any one of claims 7-14, and

- b) at least one of a reagent or a device for detecting the antibody.
- 16. An assay for the detection of a particular GPCR in a sample, comprising:
- a) providing an isolated antigenic peptide according to any one of claims 1-5,
- b) contacting the isolated antigenic peptide with the sample under conditions suitable and for a time sufficient for the antigenic peptide to bind to one or more antibodies specific for the particular GPCR present in the sample, to provide an antibody-bound antigenic peptide, and
- c) detecting the antibody-bound antigenic peptide, and therefrom determining whether the sample contains the particular GPCR.
- 10 17. The assay of claim 16 further comprising the step of binding the isolated antigenic peptide or the antibody to a solid substrate.
 - 18. The assay of claim 16 or 17 wherein the sample is an unpurified sample.
 - 19. The assay of any one of claims 15-18 further comprising, prior to the contacting, obtaining the sample from a human being.

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- 20. The assay of any one of claims 15-19 wherein the assay is selected from the group consisting of a countercurrent immuno-electrophoresis (CIEP) assay, a radioimmunoassay, a radioimmunoprecipitation, an enzyme-linked immuno-sorbent assay (ELISA), a dot blot assay, an inhibition or competition assay, a sandwich assay, an immunostick (dip-stick) assays, a simultaneous assay, an immunochromatographic assay, an immunofiltration assay, a latex bead agglutination assay, an immunofluorescent assay, a biosensor assay, and a low-light detection assay.
- 21. An isolated nucleic acid molecule encoding an antigenic peptide according to any one of SEQ ID NOS, 692-2292.
- 22. The isolated nucleic acid molecule according to claim 21 wherein the molecule encodes a naturally occurring human antigenic peptide.
 - 23. An isolated nucleic acid molecule encoding an antigenic peptide that is at least about 90% identical to any one of the antigenic peptides set forth in SEQ ID NOS. 692-2292.
 - 24. The isolated nucleic acid molecule according to claim 23 wherein the antigenic peptide is at least about 95% identical to the antigenic peptide.
- 25. The isolated nucleic acid molecule according to claim 23 or 24 wherein the molecule encodes a naturally occurring human antigenic peptide.

26. A process for producing an isolated polynucleotide comprising hybridizing a nucleotide encoding an antigenic peptide according to any one of SEQ ID NOS. 692-2292 to genomic DNA under highly stringent conditions and isolating the polynucleotide detected with the nucleotide.

- 27. A method of identifying an amino acid sequence for an antigenic peptide from a candidate polypeptide sequence wherein the antigenic peptide has a length of about 5 to about 100 amino acids, the method comprising:
- a) searching the candidate polypeptide sequence using a comparison window of the length, and
- b) selecting against amino acid sequences of the length and having at least 3 characteristics selected from the group consisting of 1) at least two consecutive prolines, 2) at least two consecutive serines, 3) at least two consecutive lysines, 4) at least two consecutive arginines, 5) at least two consecutive aspartic acids, 6) at least two consecutive glutamic acids, 7) methionine, 8) tryptophan, and 9) at least five consecutive amino acids comprising no charged amino acids.
 - 28. The method of claim 27 wherein the method further comprises selecting against at least 5 of the characteristics.
 - 29. The method of claim 27 wherein the method further comprises selecting against at least 7 of the characteristics.
 - 30. The method of claim 27 wherein the method further comprises selecting against the 9 characteristics.

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- 31. The method of any one of claims 27-30 wherein the method further comprises:
- c) selecting against amino acid sequences of the length and having at least one of the following additional characteristics 1) sequences having at least 5 consecutive amino acids that are identical to an alternative amino acid sequence from an alternative polypeptide that is different from the candidate polypeptide, 2) posttranslational modification sites, and 3) highly hydrophobic sequences.
- 32. The method of claim 31 wherein the posttranslational modification sites are phosphorylation or glycosylation sites.
- 30 33. The method of claim 31 or 32 wherein the method further comprises selecting against at least 2 of the additional characteristics.

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34. The method of claim 31 or 32 wherein the method further comprises selecting against the 3 additional characteristics.

- 35. The method of any one of claims 27-34 wherein the method further comprises performing a BLAST-type or a FAST-type analyses for the candidate polypeptide sequence.
- 5 36. The method of any one of claims 27-34 wherein the method further comprises performing a BLAST analysis for the candidate polypeptide sequence.
 - 37. The method of any one of claims 27-36 wherein the antigenic peptide has a length from 6 amino acids to about 50 amino acids.
- 38. The method of any one of claims 27-36 wherein the antigenic peptide has a length from 6 amino acids to about 20 amino acids.
 - 39. The method of any one of claims 27-36 wherein the antigenic peptide has a length of about 20 amino acids.
 - 40. The method of any one of claims 27-39 wherein the polypeptide is a protein.
- 41. The method of any one of claims 27-40 wherein the polypeptide is a human 15 protein.
 - 42. The method of any one of claims 27-41 wherein the polypeptide is a naturally occurring protein.
 - 43. An isolated antigenic peptide that is specific for the candidate polypeptide of any one of claims 27-42 that is produced according to the method of any one of claims 27-42.
 - 44. An antigenic peptide that is at least about 90% identical to the isolated antigenic peptide of claim 43.

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- 45. An isolated antigenic peptide that is an analog of the isolated antigenic peptide of claim 43.
- 46. An isolated antigenic peptide comprising a short antigenic amino acid sequence that is identical to at least 5 consecutive amino acids of the isolated antigenic peptide of claim 43.
 - 47. An isolated antigenic peptide comprising a short antigenic amino acid sequence that is identical to or contains no more than one conservative amino acid substitution over at least 7 consecutive amino acids of the isolated antigenic peptide of claim 43.
 - 48. A kit for the detection of antibodies against the candidate polypeptide of any one of claims 43-47 in a sample comprising:

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a) an isolated antigenic peptide according to any one of claims 43-47 and derived from the candidate polypeptide, and

- b) at least one of a reagent or a device for detecting the antibodies.
- 49. An isolated antibody specific for a candidate polypeptide comprising an amino acid sequence that is identical to the amino acid sequence of the isolated antigenic peptide of claim 43, wherein the antibody was produced using the isolated antigenic peptide of claim 43.
- 50. An isolated antibody specific for a candidate polypeptide comprising an amino acid sequence that is identical to the amino acid sequence of the isolated antigenic peptide of claim 44, wherein the antibody was produced using the isolated antigenic peptide of claim 44.
- 51. An isolated antibody specific for a candidate polypeptide comprising an amino acid sequence that is identical to the amino acid sequence of the isolated antigenic peptide of claim 45, wherein the antibody was produced using the isolated antigenic peptide of claim 45.

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- 52. An isolated antibody specific for a candidate polypeptide comprising an amino acid sequence that is identical to the amino acid sequence of the isolated antigenic peptide of claim 46, wherein the antibody was produced using the isolated antigenic peptide of claim 46.
- 53. An isolated antibody specific for a candidate polypeptide comprising an amino acid sequence that is identical to the amino acid sequence of the isolated antigenic peptide of claim 47, wherein the antibody was produced using the isolated antigenic peptide of claim 47.
- 54. The isolated antibody of any one of claims 49-53 wherein the antibody has high specificity and high affinity for the candidate polypeptide.
 - 55. A kit for the detection of antibodies against the candidate polypeptide of any one of claims 43-47 comprising:
 - a) an isolated antibody according to any one of claims 49-53, and
 - b) at least one of a reagent or a device for detecting the antibody.
 - 56. An assay for the detection of a candidate polypeptide in a sample, comprising:
 - a) providing an isolated antigenic peptide according to any one of claims 43-47,
 - b) contacting the isolated antigenic peptide with the sample under conditions suitable and for a time sufficient for the antigenic peptide to bind to one or more antibodies specific for the candidate polypeptide present in the sample, to provide an antibody-bound antigenic peptide, and
 - c) detecting the antibody-bound antigenic peptide, and therefrom determining whether the sample contains the candidate polypeptide.

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57. The assay of claim 56 further comprising the step of binding the isolated antigenic peptide or the antibody to a solid substrate.

- 58. The assay of claim 56 or 57 wherein the sample is an unpurified sample.
- 59. The assay of any one of claims 56-58 further comprising, prior to the contacting, obtaining the sample from a human being.
 - 60. The assay of any one of claims 56-59 wherein the assay is selected from the group consisting of a countercurrent immuno-electrophoresis (CIEP) assay, a radioimmunoassay, a radioimmunoprecipitation, an enzyme-linked immuno-sorbent assay (ELISA), a dot blot assay, an inhibition or competition assay, a sandwich assay, an immunostick (dip-stick) assays, a simultaneous assay, an immunochromatographic assay, an immunofiltration assay, a latex bead agglutination assay, an immunofluorescent assay, a biosensor assay, and a low-light detection assay.
 - 61. An isolated nucleic acid molecule encoding an antigenic peptide according to any one of claims 43-47.
- 62. The isolated nucleic acid molecule according to claim 61 wherein the molecule encodes a naturally occurring human antigenic peptide.

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- 63. An isolated nucleic acid molecule encoding an antigenic peptide that is at least about 90% identical to any one of the antigenic peptides set forth in claims 43-47.
- 64. The isolated nucleic acid molecule according to claim 63 wherein the antigenic peptide is at least about 95% identical to the antigenic peptide.
- 65. The isolated nucleic acid molecule according to claim 63 or 64 wherein the molecule encodes a naturally occurring human antigenic peptide.
- 66. A process for producing an isolated polynucleotide comprising hybridizing a nucleotide encoding an antigenic peptide according to any one of claims 43-47 to genomic DNA under highly stringent conditions and isolating the polynucleotide detected with the nucleotide.

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toticaace caaagittaa agaagactgg aagitactga agegaegtgt taccaagaaa agtggateag titeagtite cateagtage iaacaataaa attagaggoc tgagtoaaca ctgttttgat ggaciagata acctggagac cttagacttg agttataata acttggggga taacctaga tgtaagtttc aatgaattaa cttcctttcc tacggaagge ccgaatgggc taaatcaact gaaacttgtg ggcaacttca gacaggtac aaagataagc agcataccta ataatttgtg tcaagaacaa aagatgotta ggactttgga cttgtcttac aataatataa iaggectgat atetetaagg attetagate tgagtagaaa eetgataeat gaaatteaca gtagagettt tgecaeaett gggecaataa toctaactt ttottgatgo tøtgtootgg ggoagattog otgaatttgg catttggtgg gaaactggoa gtggotgoaa agtagotggg gettacaate taccaagagt taaagaetga aetaetgtgt gigtaacegt ttececegte aaccaaaate agigttata gagigaacce gaaccticc aagtittaat ggitgccatg ctctggaaga aattictita cagcgtaatc aaatclacca aataaaggaa ggcaccttic scagcaaatg teacaagcae tettgaaaat gaagaacata gteaaataat tateeattgt acacetteaa eaggtgettt taageeetgt icaggogotg accotggoto toaacaagat otcaagoato cotgactitg cattaccaa cotticaago otggtagtto igoatotica taaactcac tagcattttt attaatggcc gitatctaca ctaagctata ctgcaacttg gaaaaagagg acctctcaga aaactcacaa gicattitica aagaacaggi gcctaaatta taaattggtg aaaaatgcaa tgtccaagca atgtatgatc tgtttgaaac aaatatatga ategebaaga geaateatet eaaacagite egggitgetg ecettiegge titectaggt getacagtag eaggetgitt teceetitie agotgogaa togttictti taacaaagoc agtatcatgo aaacacttga taaaatcaca cagotgtoot goattggoag tggottottg atagagggg aatatterge ateaecectt tgtttgeeat ttectacagg tgaaaegeea teattaggat teaetgtaae gttagtgeta lgcigaaaga agcottagca gcaaaagact ttgttaacct caggicttta toggiaccat atgcitatca gigcigigca ttttggggtt attotoato titoatotgg gaagoaotto tgiaatoaot gootggtgto aottagaaga aggagaggg goagtitatt totoaaacoa tictigcag tttictecte agaaagigee atattittat taatgetage aactgtegaa agaagettat etgeaaaga tataatgaaa tegetetae taatattite eaattigete ggateteace tageaatage ttegattata tagaaagtaa aeteteggtea ataettgeal taattagac gaaacgggga gtaattatga cacgaagtac ttatgtttat ttcttagtga gctggattat cttgaacctg tgctattaar uttecteag getattaaag ecegtectag ecttaaagag etaggattte atagtaatte tattteigtt ateeetgatg gageatttga aactactaa ctaatgiggg ggittaatag tatcigaggg attiggiggc ticatgiaat gtictcatta atgaatacti cctaatatog ggaaattic catacatcit ccccatacta ttittiataa aagagectat teaatagcic agaggtigaa ctetggttaa acaagataa cttgaaaagg atcttaggtg tagtagagca atataatgtt agttttttct gatccataag aagcaaattt atacctattt gtgtattaag refecaatet etateagece egaaataatg aagtetgtta etetgatatt titteeatig eetgetigee tgaateeagt eetgtatgit iatcagtaat titticitaa gigittigig attacactac tagaaaaaaa gtaaaaggct aattgctgig igggttiagt cgattiggct agaagatgt tittaaaaca atattaacag ctgttaggtt aaaaaaatag ctggacattt gtittcagtc attatacatt gctitggtcc ggiaatcca ctcttaagaa ctatacattt gtatgalaat cctctgtctt ttgtggggaa ctcagcatct cacaatttat ctgatcttca cattigeat citgracate aetgectieg tecaaatigt trataggeit gattietgig telaaettat teatgggaat etataetgge acataggea ttacittatt atgitticae ttgecatect tgacataaga gaactataaa ttttgittaa geaatttata aatetaaaae ctagcatga ttaagcatgt cgcttggcta atcitcacca attgcatcit ttictgccct gtggcgtttt tticattigc accattgatc sacaagataa agaacagcig ttaatattt ttaaaaatct attitaaaat gtgatttict ataactgaag aaaatatcti gcraatttia patatitac teggaagotg gatgattogt ettactetet getteattit ettegtteca trattitica acctectigt tattitaaca igacicita igcaaattia aacacagaag ataacagcci ccaggaccac agigiggcac aggagaaagg tacigcigat xeaaagacet gagggetaet ggteegaetg tggeacaeag teggeecaet etgattatge agatgaagaa gatteetttg etteagtiae ggeatetgig geiggatgae aacagetiga eggaggigee igigeaeeee eteageaate igeecaeeet cetaatgttt catecttaat etcaggacaa ettaetgeag ggecaaaaaa gggaetgtee cagetagaae tgtgagagta ctcagacag ttctgaccag gtgcaggcct gtggacgagc ctgcttctac cagagtagag gattcccttt ggtgcgctai naggiggit gtotggaaca ggatttctac tacgactgig gcatgtactc acattigcag ggcaacctga cigitigcga tecetagte attegtggtg caagcatggt geageagtte eccaatetta caggaactgt ceaectggaa agtetgaett

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MPGPLGLLCF LALGLLGSAG PSGAAPPLCA APCSCDGDRR VDCSGKGLTA

fatgaaatac aatattgtac tcagtgtttt gaattattaa agtttctaga aagcaaaaa a

LSGLKELKVI, TLQNNQLKTV PSEARGLSA LQSLRLDANH ITSVPEDSFE SLVQLRHLWL DDNSLTEVPV HPLSNLPTLQ ALTLALNKIS SPDFAFTNL

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KELGFHSNSI SVIPDGAFDG NPLLRTIHLY DNPLSFVGNS ASHNLSDLHS
LVIRGASMVQ QFPNLTGTVH LESLTLTGTK ISSIPNNLCQ EQKMLRTLDL
SYNNRDLPS FNGCHALEEI SLQRNQIYQI KEGTFQGLIS LRILDLSRNL IHEIHSRAFA
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SLSVPYAYQC CAFWGCDSYA NLNTEDNSLQ DHSVAQEKGT ADAANVTSTL
ENEEHSQIII HCTPSTGAFK PCEYLLGSWM IRLTVWFIFL VALFPNLLVI LTFASCTSL
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AGFLAVFSSE SAIFILMLAT VERSLSAKDI MKNGKSNHLK QFRVAALSAF
LGATVAGCFP LFHRGEYSAS PLCLPFTGE TPSLGFTVTL VLLNSLAFLL
MAVYTKLYC NLEKEDLSEN SQSSMIKHVA WLIFTNCIFF CPVAFFSFAP LITAISISPE
IMKSVTLIFF PLPACLNPVL YVFFNPKFKE DWKLLKRRVT KKSGSVSVSI
SSQGGCLEQD FYYDCGMYSH LQGNLTVCDC CESFLLTKPV SCKHLIKSHS
CPALAVASCQ RPEGYWSDCG TQSAHSDYAD EEDSFVSDSS DQVQACGRAC
FYQSRGFPLV RYAYNLPRVK D

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gettegecec caacaactte gegeteetg egeacategt gageegeetg tictaeggea agagetacta ceaegtgtae aaggeteactg eggeteggg accegttigt taitacttt gegteeggg agitecaggt gegeteggg gaatteaggt gegeteggg gaatteaggt gegeteggg gaatteaggt gegeteggg gaatteaggg gaaggtegggggggggg	integras a subsessible transmotes generated englighted and gogge a controlled and gogge the granting a confection of the gogge and gogge and gogge controlled and gogge gogge and gogg and g	MQVPNSTGPD.NATLQMLRNP AIAVALPVVY SLVAAVSIPG NLFSLWVLCR RMGPRSPSVI FMINLSVTDL MLASVLPFQI YYHCNRHHWV FGVLLCNVVT VAFYANMYSS ILTMTCISVE RFLGVLYPES SKRWRRRRYA VAACAGTWLL LLTALSPLAR TDLTYPVHAL GIITCFDVLK WTMLPSVAMW AVFLFIFIL LFLIPFVITV ACYTATILKL LRTEEAHGRE ÓRRRAVGLAA VVLLAFVTCF APNNFVLLAH IVSRLFYGKS YYHVYKLTLC LSCLNNCLDP FVYYFASREF QLRLREYLGC RRVPRDTLOT RRESLFSART TSVRSFAGAH PFGMFGATRP GLOROFSVF	gaatteggee aaagaggeet atgettetet gaagactige ageaaggett getgaggete eagaagata geeccagtgt titggagtgg tittgaatgt gattetgaga teagactige ageaaggett etegaggete acagaagata geeccaa cettggagte titggagastit titggaatgt tittgaatgt gaattetget tagetggaga teetggette tageccagtt ataaagatet ggaactatt teectcaaga tgacaaacag titegtiette tgeccagttt ataaagatet ggaactattit titattagt titeettgit ggaattatit gaagtitgit tgeaacctgg gettitatae agaagaatae gaateacaagg tigtgragea tetaettaat taattgett acagecgatt teetgetae tetageataa agaagaaaa tigttgitga ettaggtgg geaectitga agetgaagat attecatige caagtagae agateacag eetgecatt gaateaagaa attecatige caagtagae agateacea agateacaa agaataatt caataatet eetagaata tacaaccgt gtgggeaaa gaattggaa agaattgga tigtaggaaaa tittaaaaagg aatttggaa aaattggga tigtagaaaa teetaaaaga aattggga tigtaggaata teetaaaaga aaattggaa tigtetgaaaa attecaaaa tigtaaaaagg eteteaaaaaga taaccaaaa tigtaaaaagg eteteataaa catactitta gtgaccaagga tittaatatg eetaaateaa agettagt eetaaaaaga attaccaaaa tigtaaaaagga eteteaaaaagaagateat aactgattggt teaaccagga titecateta tagettigtt eetaaccaaa tigtegaaa etgtegtgaaa etgteetaaaaagaaaagaa	gctcagaaag aaaaataag atgtgaaaa aatgcataaa agacaggat tittgtgcta ccaattctgg cctactgga ccataaagtt aattatagct tigaaagata aaaaaaaaa aaaagcggcc gc MTNSSFFCPV YKDLEPFTYF FYLVFLVGII GSCFATWAFI QKNTNHRCVS IYLINLLTAD FLLTLALPVK IVVDLGVAPW KLKIFHCQVT ACLIYINMYL SIIFLAFVSI DRCLQLTHSC KIYRIQEPGF AKMISTVVWL MVLLIMVPNM MIPIKDIKEK
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		LR80	NM_013308	NP_037440.1
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		LS160435 Receptor	Platelet Activating Receptor Homolog (H963)	Platelet Activating Receptor
		160435	160889	160889
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Protein A

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DNENYPNVKK ALINILLVIT GYIICFVPYH IVRIPYTLSQ TEVITDCSTR ISLFKAKEAT LLAVSNI, CF. DPILYYHLSK AFRSKVTETF ASPKETKAQK EKLRCENNA INVGCMEFKK EFGRNWHILT NFICVAIFLN FSAILLISNC LVIRQLYRNK

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161024 Protein A

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GKRRSSLDGS ESAKTSLQVT NLVSAIVFLY DSLTGVPILV VSFFSLKSDS APPWMVLAVL WCSMAQTLLL PSFIWSCERY RADVRTVWEQ CVAIMSEEDG DDDGGCDDYA EGRVCKVRFD ANGATGPGSR DPAQVKLLPG RHMLFPPLER VHYLQVPLSR RLSHDETNIF STPREPGSFL HKWSSSDDIR VLPAQSRALG GPPEYLGQRH RLEDEEDEEB AEGGGLASLR QFLESGVLGS GGGPPRGFF FREEITIFID ETPLPSPTAS PGHSPRRPRP LGLSPRRLSL GSPESRAVGL PLGLSAGRRC ŠLTGGEESAR AWGGSWGPGN PIFPQLTL	toccaggige cegicigatg gggagatgge tgatgeccag aacatticac iggacagece agggagtgig ggggectggg cagtgectgt ggicitigec chaitetiec igcigggeae agtgggeaat gggelggige iggeagtget eetgeaged ggeoegagtg eetggeagga geetggeage accaeggaee tgiteateet caacetggeg giggetgaee tegetteat	cotgigatgo gigacottoc aggacaccat ciacacgotg gatgoctggo totttggggo octogtotgo aaggacgigo acotgotcat ciacotcaco atgiacgoca gcagotttac gotggotgot giotocgigg acaggiacot ggooglgogg cacocgotgo gotogogogo octgogcacg cogogtaacg ocogogcogo agtggggotg gigtggotgo tggoggogot	cttotoggog coctacotca gotactaogg cacoglgogo taoggogogo tggaggototg ogtgocogoo tgggaggaegacg cgegocogoo tgggaggaegacg cgegocogoo gogocogoo tgggaggaega cgegocogoo gegocogoo gggocacot tegotgocogg ctaocogogo cocogogogo gocogogogo ggcogogogo gocogogogo gocogogogo gocogogogo gocogogogo gocotgotogo gocotgotogo gocotgotogo gototaocogo cotogotogog gocogoacoa egegotocatco cigigottot	gglacggcog citegentic ageneggoea octangenig cogentigges teanantgen tggentange caantenge caantenge caantenge citegenia generange cannot cogenegent octanange in the cogenegent octanange categoegang cogenerate generange specification octanange cogenerate generate gen	MADAQNISLD SPGSVGAVAV PVVFALIFLL GTVGNGLVLA VLLQPGPSAW QEPGSTTDLF ILNLAVADLC FILCCVPFQA TIYTLDAWLF GALVCKAVHL LIYLTMYASS FTLAAVSVDR YLAVRHPLRS RALRTPRNAR AAVGLVWLLA ALFSAPYLSY YGTVRYGALE LCVPAWEDAR RRALDVATFA AGYLLPVAVV SLAYGRTLRF LWAAVGPAGA AAAEARRRAT GRAGRAMLAV AALYALCWGP HHALILCFWY GRFAFSPATY ACRLASHCLA YANSCLNPLV YALASRHFRA RFRLWPCGR RRRHRARRAL RRVRPASSGP PGCPGDARPS GRLLAGGGQG PFPDFGDVHG GFA A DGDF	algregatiga coccegagacaco taccigago tegoregocac oggoagotot gigoregago egorigageg coccagegago coccagegago coccaacegos acoticaaca goticotigago cagocogaco gagocoagot cocigaagga cotigatigaco acegegacoca tigagagotot gotigtegaco ategagos egorigana gagocoagot tacacegotag tegatoacota cogotocota
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	Galanin Receptor GalR3				Galanin Receptor GalR3	Urotensin-II Receptor (GPR14)
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cigggotigo itotigocot ictigacigig gcagotigto goccagiaco accagiacoc gciggogog eggacggogo gcalogiaco gcalogiaco gcalogiaco gcalogiaco gcalogiaco accocitoc ictacacgot gcicaccagg aactacogog accaccigog eggocogogiacogogogogogogogogogogogogogogogogogogog	CALUESTICA EBUCGEACH ENGINEER BICCHERERS MALTPESPSS FPGLAATGSS VPEPPGGPNA TLNSSWASPT EPSSLEDLVA TGTIGTLLSA MGVVGVVGNA YTLVYTCRSL'RAVASMYVYV VNLALADLLY LLSPFIVAT YVTKEWHFGD VGCRVLFGLD FLTMHASIFT LTVMSSERYA AVLRPLDTVQ RPKGYRKLLA LGTWLLALL TLPVMLAMRL VRRGPKSLCL	FAWGFKAHKA YLILLFATSI AGPGLLIGLL YARLARAYRR SQRASFKRAR RPGARALRLV LGIVLLFWAC FLPFWLWQLL AQYHQAPLAP RTARIVNYLT TCLTYGNSCA NPFLYTILTR NYRDHLRGRV RGPGSGGGRG PVPSLQPRAR FQRCSGRSLS SCSPOPTDSL VLAPAAPARP APEGPRAPA	algectigea algecagige geocaggggg cactitigace cigaggacti gaacetgat gaegaggcae tgagacteaa giacetgggg cocageagg cactitigace cigaggacti gaacetggg geogetgtigg gaegetgtigg cactiggggggggggggggggggggggggggggggggggg	CETTEGICE (EGCATITIC) gracegotard gritgagalg grotgocygg corcayigot caacegoard geocygagog tegaacegor tegaacegor atgreegog agreetiggg geocygaacegora tegaacegora tegaacegora tegaacegora tegaacegora tegaacegora egocygaaceg geocygaacegora geocygaacegora egocygaacegora geocygaacegora geocygaacegora tegaacegora tegaacegora egocygaacegora geocygaacegora geocygaacegora geocygaacegora tegaacegora	CIBCICATEC ABGABECCAA ABBCAGEEGE (tripcagcag ccaggiccag atacaccigc aggetocage agcacgateg BEBCCEGAGE aggetocage agcacgateg BEBCCEGAGE aggreeges agagetit tripcagcage gegocegate gegocegate aggetatit grantifies aggetotit geacgicage transparation aggetotit gegociegit cacagegate transparation aggetotit gegociegit geaggetot geacagetot geocagate transparation aggetotit gegociegit geocagace effectitata geocicatit cagcocgit cagagage colgination aggetotic gegociegit geoalogici cagaococge cacagetoc acagetoca acageaga accititiga tetigggetoc cigggeaget gegocacoc colgidage aacgatege cagagagaco gatocatect ga	MACNGSAARG HEDPEDLNLT DEALRLKYLG PQQTELFMPI CATYLLFVV GAVGNGLTCL VILRHKAMRT PTNYYLFSLA VSDLLVLVG LPLELYEMWH NYPFLLGVGG CYFRTLLFEM VCLASVLNVT ALSVERYVAV VHPLQARSMV TRAHVRRVLG AVWGLAMLCS LPNTSLHGIR QLHVPCRGPV PDSAVCMLVR PRALYNMVVQ TTALLFFCLP MAMSVLYLL IGLRLRRERL LLMQEAKGRG SAARSRYTC RLQQHDRGRR QVTKMLFVLV VVFGICWAPF HADRVMWSVV SQWTDGLHLA FQHVHVISGI FFYLGSAARP VLYSLMSSRF RETFQEALCL GACCHRI RPR HSSHSI SRMT TGSTI CDVGS I GSWVHPI AG NINGPEAOCHT DES	atggetaace ttgacaaata cactgaaaca tteaagatgg gtagcaacag taccagcact getgagattt aetgaatgt cactaatgtg aaattteaat acteceteta tgeaaccace tataicetea tatteattee tggtettetg getaacagtg cagcettgtg ggttettgtge egetteatea geaagaaaa taaagecate attteatga teaacetete tgtggetgae ettgeteatg tattatettt
	NP_061822.1		NM_006056			NP_006047.1	NM_014499
	Urotensin-II Receptor (GPR14)		G Protein- Coupled Receptor GPR66			G Protein- Coupled Receptor GPR66	Purnergic Receptor P2Y10
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	238		239			540	541

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					NP_055314.1						NP_042597.1	÷					٠		MM_006679		
						Receptor P2Y10					G Protein-	Coupled Receptor	Ls161293	[Herpes virus]			-		177147 Neuromedin K	Receptor-Like	(NK4R)
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ttaaatatat taaaaatcat atgaaaaat MASPAGNLSA WPGWGWPPA ALRNLTSSPA PTASPSPAPS WTPSPRPGPA HPFLQPPWAV ALWSLAYGAV VAVAVLGNLV VIWIVLAHKR MRTVTNSFLV NLAFADAAMA ALNALVNFIY ALHGEWYFGA NYCRFQNFFP ITAVFASIYS MTAIAVDRYM AIIDPLKPRL SATATRIVIG SIWILAFLLA FPQCLYSKIK VMPGRTLCYV QWPEGSRQHF TYHMIVIVLV YCFPLLIMGI TYTIVGITLW GGEIPGDTCD KYQEQLKAKR KVVKMMIIVV VTFAICWLPY HIYFILTAIY	QOLNRWKYIQ QVYLASFWLA MSSTMYNPII YCCLNKRFRA GFKRAFRWCP FIHVSSYDEL ELKATRLHPM RQSSLYTVTR MESMSVVFDS NDGDSARSSH QKRGTTRDVG SNVCSRRNSK STSTTASFVS SSHMSVEEGS	atggatgaa caggaaatct gacagtatct tetgecacat gecatgacac tattgatgac tteegeaate aagtgtatte cacettgtac tedatgatct ctgtgatgatet etgettgat tedatgatct etgtgage aatggettg tgetetatgt ecleataaa acetateaca agaagteage ettecaagta tacatgatta atttageagt agcagateta etttgtgtgt geacactgoc teteogtgtg gtetattatg tteacaaagg catttggete	ittegrigaci tettigigeeg ecteageaec targentigi atgicaaeet etatigiage ateitetta tgacagecat gagettitie eggigeatti tecagteeag aacattaatt tggtacae gaaaaagee aggittegigt grgtaggiat ttggattiti grgattitiga ecagitetee atticiaatg gecaaaceae aaaaagatga gaaaaataat accaagtget tigageeece acaagaacaat caaactaaaa ateatgitti ggtetteeat targgieai tgttigtigg etitateate cettitigtia ttataattgi	ctgitacaca atgaicatti tgaccitact aaaaaaatea atgaaaaaa arcigicaag icataaaaag gcialaggaa tgaicatggi cgitacaca atgacogit goctititag icagaticat gccataicat atteaacgia ccattcacci tcattitita cacaatgaaa ctaaaccig tgaticigic citagaatgc agaagtccgi ggicataacc tigtcicigg ctgcatccaa tigtigciti gacccictcc tatatiticit ticigggggi aacittagga aaaggcigic tacattcaga aagcattcit tgiccagcgi gacttatgia cccagaaaga aggccictit eccagaaaa aateraaaat ataa	MDETGNLTVS SATCHDTIDD FRNQVYSTLY SMISVVGFFG NGFVLYVLIK TYHKKSAFQV YMINLAVADL LCVCTLPLRV VYYVHKGIWL FGDFLCRLST YALYVNLYCS IFFMTAMSFF RCIAIVFPVQ NINLVTQKKA RFVCVGIWIF VILTSSPFLM AKPQKDEKNN TKCFEPPQDN QTKNHVLVLH YVSLFVGFII PFVIIIVCYT MILTLKKS MKKNLSSHKK AIGMINVVTA AFLVSFMPYH IORTIHLHFL	HNETKPCDSV LRMOKSVVIT LSLAASNCCF DPLLYFFSGG NFRKRLSTFR KHSLSSVTYV PRKKASLPEK GEEICKV ccacgcgtc gccggcigca cggtcgcacc ggcagcggc caggctcgg ctctctccc gctgcagcag ccgcgtgcc ggcccactg ggctcggatc cggccccggc ccctcggca ccgctgctc tggccccgg cccggccccg cggaccatgc gctggcgcc ccaggggaa acccgacccg gccaagggc cgcaaagacg aggctcccgg gccgggcc ctccggccc gccagctctc ggccggcgc ctgcccgcg tcccggagc gcgtgagcct gcggggccat ggagcgcgc ctccggccg ccagctctc ggccgcgcc ctgcccgcg tcccggagc gcgtgagcct gcggggccat ggagcgcgc ctccggccg ggcgttgaa cgcttcgggg gcgctggcgg gcgatgcgg gggggcgc ggtgggcgc ggtgggcg ggcgctgat cgctcggcgc atcgggcca cggtgagcc cacgcgcgg gtctcggc agcttgggc gcggtgctgg ccgcgctcat ggagcaa cggtggcgg caacgcgctg gtcatgctc cttcgtggc
NP_006670.1		NM_006639			NP_006630.1	NM_007232
Neuromedin K Receptor-Like (NK-4R)		Cysteinyl Leukotriene CYSLT1	Keceptor		Cysteinyl Leukotriene CYSLT1 Receptor	Histamine H3 Receptor
177147		177168			177168	177191
545	-	546			547	548

ataccegec cagcagegg acacecege geaggegg agagtecte tegtggggg tegtggggg cottc cegcigace gaccagocat cotgagctgg gagtacctgt cegegggcag ctocatecce gagggccact getatgcega gttettetac cgactogage etcogracce agaacaactt etteotgete aacetogeca totoegaett cotogtogge geettetgea teccactgta tgdaccetae gtgetgacag geogetggae ettoggeogg ggeetetgea agetgtgget ggtagtggae taccigetg geactetge acactete geottecae ateggeete taccigetgt geacetect tgcotteaae ateggeete teagetaega cogetteetg teggteacce gageggetete

Homo sapiens

sapiens

GNALVMLAFV ADSSLRTQNN FFLLNLAISD FLVGAFCIPL YVPYVLTGRW

NP_009163.1

Histamine H3

177191

Receptor

collected etterataag celeagect gecettica ecettetic caccaactet etergeece aaaagtgica aggggeecta ocaccitic gragitarig gitiggigite ticceaaage aageaectigg gigigeicea ggetteetige ectageagit tgeetetigea ggaacctoga agotgitoto igottitoca itotgggigi ittoagaaag atgaagaaga aaacaigiot gigaactiga igitogiggg aactegtact tecteateae ggettecaee etggagttet ttaegecett ecteagegte aeettettta aeeteageat etaeetgaae gt gagggg c gaaggcgct gaggccgggg aggcgacoct cgggggggg ggtggc ggtggggggg gctocgtggc ttcacocaco atica gagge gracocgoot coggot ggg ggg etegag aggoagoegg coocgagcoc cotoocgagg coagoote accaccca cogoctegot gotegegotig ctegoagaag gegoacegeg aggocatgoo gotecacag tategegeteg caaggegtge aggggeggte cagaggaggt geoegggeag gggeegette gecatgtget gtgeaceegt gecaegege iccagcicog gcagcicoto gaggggcact gagaggcogo gotoactoaa gaggggctoc aagcogtogg ogtoctoggo Beagceac coteccateg aggogcotto otgegitiggo cagagggooc otcaciggot ggaotggagg otgggiggo ttgccceggc cactctgttt gctcacccag gacctctggg ggttgttggg aggaggggc coggctgggc ccgagggtcc egigeacaca cotgeacace cotgeacaca cotgeacace grecotote coggacaage coaggacact gootttgetg gotocotgga goactgotgg aagtgagtgg cocaccagag cotocotcag coacgootot otcagoocag gtotootggg gecotgecc eceacatict ggetecaccg gggagggaca gtetggaggt eceagacatg etgeocaece eetgetggt -Bedaagget teeggetgag etgtgecage tgettetgee caeceegeet etgggeteae aecageeetg gtggecaage ctogotegag aagogoatga agatggtgto coagagotto accoagogot ttoggotgto togggacagg aaagtggoo daccototg tgocaccaca gottocgoog ggoottoaco aagotgotot goooccagaa gotoaaaato cagooocaca altigged tectgodde tacceggote gtteccodag gggtgagod egoegtgtet gtggeddet ettaatgod agicgcigge egicategig ageatettig ggetetgetg ggeeceatae aegetgetga tgateateeg ggeegeetge itgittaato aagagagaca aaattgotga ggagotoagg gotggattgg caggtgtggg otoocaogoo otootocolo MERAPPDGPL NASGALAGDA AAAGGARGFS AAWTAVLAAL MALLIVATVL categocact gegrecinga ctactegrac gaaacciect tetegoticit greggocaac teggotgrea accitered otgeatgote etetgeetgt geoogetgeg etgeeetgea aacegtgagg teacaataaa gtgrattitt ttaaaaaaa aaaaaaaa aaaaaaaa

NM 020155

Coupled Receptor

G Protein-

getetgeete acetegetee aggtgeaceg ggtggeaege agacactgee agegeatgga cacegteace atgaaggege

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₩.	Homo	Homo		Homo	Homo
	<u>р</u> ,	⋖		ρ ₄	∢
gegocaacog celggggcce ttgocottet ggettetera etgetgccce gtetgeetge agttetteae ettgaegett atgaacetet actttgecca ggtggtgtte aaggecaagg tgaagegteg geoggagag ageegagge ttgetegetgt cegaggggcc tttgtggggg cetegetget etttetgetg gtgaaegtge tgtgtgetgt getetecat eggegegeae agocotggge cetggetget gtgaaegtge tgtgtgetgt getetecat eggegegeae agocotggge cetggetget gtgaaegtge gtgatetgt getetecat eggegegee eteggeggg eggeggggggggggggg	MESNI.SGLVP AAGLVPALPP AVTLGLTAAY TILYALLFFS VYAQLWLVLL YGHKRLSYQT VFLALCLLWA ALRTTLFSFY FRDTPRANRL GPLPFWLLYC CPVCLQFFTL TLMNLYFAQV VFKAKVKRRP EMSRGLLAVR GAFVGASLLF LLVNVLCAVL SHRRAQPWAL LLVRVLVSDS LFVICALSLA ACLCLVASGR	ctrotttaaa titotticta ggatgitcae trottorca caatgaalga gtgrcactat gacaagcaca tggactittt tiataalagg agcaacactg atactgtoga tgactgaca ggaacaaagc tigtgattg titgigtgt gggacgitt tetgcctgt tattititt totaaticte tggicatege ggcagtgate aaaacagaa aatticatit eccettciae tacetgtigg ciaattiage tgetgccgat tettegetg gaattgccta tgattectg atgittaaca caggcccagt ticaaaaact tigactgtca accgctggtt tetcogtcag gggcottetgg acagtactic etcaccaact tgetggttat egcettggag aggcacatgt caatcatgag gatgegggic catagcaaca tgactgrac etgetgaac etgetgica etgetgaga aggcacatgt caatcatgag	gggoggtoco cacacigggo tggaattgoc totgcaacat ctotgcotgo tottocotgg cococattra cagcaggagt facettgttt totggacagt grocaacte totacagga tracttgttt totggacagt grocaacte totacagga totacagga totacagga accaacga grocaacte tgtotcogca incaagtggg tocatcaggo googgaggac acccatgaag ctaatgaaga cggtgatgac tgtottaggg gogtttgtgg tatgotggac occgggoctg gtggttctgc toctogacgg cotgaactgc aggcagtgtg gogtgcagca tgtgaaaagg tggttoctgc igotggogct gotcaactoc gtottgaacc ccatcatcta ctoctacaag gaogaggaca tgttgaaaagg tggttoctgc igotggogct gottcaactoc gtotggaacc ccatcatcta ctoctacaag gaogaggaca tgtatggaac catcatcaa gaategaca gaategaca gaategaca gaategaca gaategacaa gaategacaaa gaategacaa gaategacaaa gaategacaaaa gaategacaaaaaaaaa gaategacaaaaaaaaaaaaaaaaaaaaaaaaaaaaaaaa	aaagcactic claaactetg gatgcietic ggccacca ggtgatgact gictiagg MNECHYDKHM DFFYNRSNTD TVDDWTGTKL VIVLCVGTFF CLFIFFSNSL VIAAVIKNRK FHFPFYYLLA NLAAADFFAG IAYVELMFNT GPVSKTLTVN RWFLRQGLLD SSLTASLTNL LVIAVERHMS IMRMR VHSNL TKKRVTILLL LVWAIAJFMG AVPTLGWNCL CNISACSSLA RIYSRSYL VF WTVSNLMAFL IMVVVYLRIY VYVKRKTNVL SPHTSGSISR RRTPMKLMKT VMTVLGAFVV CWTPGLVVLL LDGLNCRQCG VQHVKRWFLL LALLNSVVNP IIYSYKDEDM VGTVMKKMACC FROFNDFRRP SRIPSYL SR STICSOVTFD SISOGAVCNK STS	alggecceg gegageget getiggeget citedata lightactigge cytiggeget citedagges cettigges entities alggested getiggeget citedagges the cytigges getigged cytigges cytigged cytigges cytigged cytigges getigged cytigges cytigges getigged cytigges cytigges getigged cytigges cytigges getigged cytigges getigged gytigges getigged gytigges
	NP_064540.1	NM_012152		NP_036284.1	AF411107
	G Protein- Coupled Receptor ORF4	Lysophosphatidic Acid Receptor Edg7		Lysophosphatidic Acid Receptor Edg7	G Protein- Coupled Receptor GPR78
	177387	180956		180956	189873
	551	552		553	554

	P Homo sapiens		A Homo sapiens		P Homo sapiens	A Homo sapiens
legeogliget egeogaectig eacoccagitg tgeggeaegg etgeeteate cageagaage ggegeogeca eegegecacc aggaagatitg geattgetat tgegaectic eteatorget tigeocogta tgreatgacc aggetiggegg agetegracc ettegracce ettegracce ettegracce aggaaegeca ecteatorget tgeocogaag tgeotgacct acageaagge ggtiggeoga cegittaagt acteticiget eegeeggeog teoggeoag eattgitgeac eggetigeta aggaaecco gegeocagca teoaccatg acagetetet ggatgitgec ggeatgitge acagetetet gaagaaacc egtecacca	caacggctct gtggacacag agaatgattc ctgcctgcag cagacacat ga MGPGEALLAG LLVMVLAVAL LSNALVLLCC AYSAELRTRA SGVLLVNLSL GHLLLAALDM PFTLLGVMRG RTPSAPGACQ VIGFLDTFLA SNAALSVAAL SADQWLAVGF PLRYAGRURP RYAGLLLGCA WGQSLAFSGA ALGCSWLGYS	SAFASCSLRL PPEPERPRFA AFTATLHAVG FVLPLAVLCL TSLQVHRVAR RHCQRMDTVT MKALALLADL HPSVRQRCLI QQKRRRHRAT RKIGIAIATF LICFAPYVMT RLAELVPFVT VNAQWGILSK CLTYSKAVAD PFTYSLLRRP FRQVLAGMVH RLLKRTPRPA STHDSSLDVA GMVHQLLKRT PRPASTHNGS VDTENDSCLQ QTH	atggaaaaac tteagaatge ttoctggate taccageaga aactagaaga tecattecag aaacacetga acageacega ggagtatetg gecttectet geggaceteg gegeagecae ttettectee cegtgfetgt ggtgtatgt ceaatititig tggtggggg cattggeat geotggtgt tetgeagea caggetatga agacgeccae caactactae etetteage tggeggtete tgaceteetg gtoctggtgt tetgeagec cegggetatga agacgeccae caactactae etetteage tggeggtete tgaceteetg gtoctgetee ttggaatgec cetggaggte tatgagatgt ggegcaacta coettiettg tteggaatge cattgaagete tatgagatgt ggegcaacta coetteetg	cagogligae godiaciping coatoctors occupied against against against a cagogligae godiaciping coatoctors occupied against agains	cttigtggag ctgaccgaag atataggicc coattocca tgroatcigc ccagcggaac atcitcciga cagaagcca cttigtggag ctgaccgaag atataggicc coattocca tgroatca ccatgcacaa ctctcacctc ccaacagcc tctctagtga acagaitca agaacaaact atcaaagct ccacttaac aaaacctga MEKLQNASWI YQQKLEDPFQ KHLNSTEEYL AFLCGPRRSH FFLPVSVVYV PIFVVGVIGN VLVCLVILQH QAMKTPTNYY LFSLAVSDLL VLLLGMPLEV YEMWRNYPFL FGPVGCYFKT ALFETVCFAS ILSITTVSVE RYVAILHPFR AKLQSTRRRA LRILGIVWGF SVLFSLPNTS IHGIKFHYFP NGSL VPGSAT CTVIKPMWTY NFIIOVTSFL FYLLPMTVIS VI YYL MAI RI KKDK SI FADF	GNANIQRPCR KSVNRMLFVL VLVFAICWAP FHIDRLFFSF VEEWSESLAA VFNLVHVVSG VFFYLSSAVN PITYNLLSRR FQAAFQNVIS SFHKQWHSQH DPQLPPAQRN IFLTECHFVE LTEDIGPQFP CQSSMHNSHL PTALSSEQMS RTNYQSFHFN KT atgctggcag ctgccttagc tccagcagca tgaatgtgtc ctttgctcac ctccactttg ccggagggta cctgccctct gattccagg actggaggaac catcatcccg gctctcttgg tggctgtctg cctggtgggca ttcgtgggaa acctgtgtgt
	CAC34041.1		NM_020167		NP_064552.1	LG94108
	G Protein- Coupled Receptor GPR78		Neuromedin U Receptor 2		Neuromedin U Receptor 2	G Protein- Coupled Receptor
: ·	189873		189874		189874	189884
•	555		556		557	558

	Homo sapiens	Homo sapiens	Homo
	Д	∢	Q .
gattggcatc ctocttoaca atgottggaa aggaaagoca tocatgatoc actooctgat totgaatoto agoottggotg atototoct cotgottgttt totgcacota tocagagota ggogatotoc aaaagtgttt gggatotagg otggtttgto tgoaagtot tocagagota tatoocaa tgoattggat gogatotaa gggatotagg otggttgto tgoaagtot totgaottgtt tatoocaa tgoattggat gacaatogt gtggtggoca aagtatgott catgatga agtgaocoag coaagaatgg tggaaatgt gottggact tggttagoca gatacocotg coggaatgg totttagoac catoaggoat catgaaggtg tggaaatgtg cotottggat gacagott tggttgga agottatga agttatatgoa agtttagta agottacoc actottggac tttagocot cattatttt tgocagottt tattotaga agottataga coaatgtaaa aaacgaggaa ctaagacta aaaatottaga aaccagata cgttaaagga agottataga accaatgtaaa aaacgaggaa ctaagacta tttagoctt cattatttt tgocagottt tattotaga gagottataga goattgaaa aaccagot cotttaga gatagottaga tgataacota cattottota gaaaatoto cattotaga gatagottaga tgataacoaa aaaacotoca actgitotag agtotcagga aacacoagot gaaaaggat tocatotoca aaaaocotoca actgitotag agtotcagga aacacoagot tgataacoaa aaaacotoca actgitotag agtotcagga aaacacoagot tocottot ggcaaaagga aaacagaaa aaacagaga aaacagaa aaaaagaga aaacagac tocottot ggcaaaagga aaactgagaa agacaatgac cattoocotg ggaacatgaa gatcaaagaa caggggaagg catgaaatag	MLAAAFADSN SSSMNVSFAH LHFAGGYLPS DSQDWRTIIP ALLVAVCLVG FVGNLCVIGI LLHNAWKGKP SMIHSLILNI. SLADLSLLLF SAPIRATAYS KSVWDLGWFV CKSSDWFIHT CMAAKSLTIV VVAKVCFMYA SDPAKQVSIH NYTIWSVLVA IWTVASLLPL PEWFFSTIRH HEGVEMCLVD VPAVAEEFMS MFGKLYPLLA FGLPLFFASF YFWRAYDQCK KRGTKTQNLR NQIRSKQVTV MLLSIAIISA LLWLPEWVAW LWVWHLKAAG PAPPQGFIAL SQVLMFSISS ANPLIFLVMS EEFREGLKGV WKWMITKKPP TVSESQETPA GNSEGLPDKV PSPESPASIP EKEKPSSPSS GKGKTEKAEI PILPDVEQFW HERDTVPSVQ DNDPIPWEHE DQETGEGV	alignatical caccatico coagicatoa gggaactott coactitiggg gagggtocot caaacoccag glocototac tgocagtggg grocipal george ggaggggot toggaatot gggocotott ottoatgoto otgotggact tgactgota gggocotot gggaact gggggact gggcgggact gggcgggat tggcaagacg cotgocotoc gaaaattigt citogtotto cacctotigot tggtggacot gdggcggc otgacoctoa tgccotigo catgotococ agocotgoco tottigacoa cacctotigot ggggagggact tgggcggc catgotococ agocotgoco tottigato caccidatiti otgagggtgt gdttigtoag cotggcatoc attigatoc agocotaca gagggaggga gdttigtoag catgotococ catgogctac gaggtgcgca tgacgctggg gctgggggc tctgtgctgg taggcgatoc agtgttiggga agggtotoct gggagggaagg agotocoagt gggggtggg gatggggaaggaaggaaggaaggaag	cigotocica facticiggi ciactgoago atgiticogag tegocogogi egotgocatg coagaceggo ogotgococoga egiggategag acacocoggo aacgotocga atloticiago agococoaga egococoaca ocogacegti egggagga aagoagat egotocoa ogateggica cagtoggga gococoaga ecacocoaca ocogacegti egggagga aagoagat egotocoa egoteggga gotocoaca ocogacegti egotatigga egotocoa egotoggagga etotitigot traciticoa ocottotito taggatego tocacegoca troacegoa egotocaga egotoca ocottotito taggatego coagocoga agocoagate troacegoa agoagatego tocaticoa agocagate ocotagotegoa egotocoa egotocoa egotocoa egotocoa egotocoa agoagatego cottoticoaga etotocoacoa ocoagococa agoagagaco acotegotat gaotticgaa tocaggocoa atag
	ENSMPRT1140 67	NM_031936	NP_114142.1
Ls189884	G Protein- Coupled Receptor (Ls189884,	G Protein- Coupled Receptor GPR61	G Protein-
	189884	189895	189895
	559	095	561

ggocototge gegeotgcot geoeggocoag accaeggtte eggatggtga geocaatgct cacaatagca aagaggatga gegecagtgg caggaagaac tecagcaggt acagtgeotg gtgecagegg agegaggeeg agggettegt geocaecetg tagetgagge aggaggggee ggagaaggtg etcaggagca ggtgeoegtt gaggagaagg atgeocaece agagteeece

sapiens	Homo sapiens		Homo sapiens	Homo sapiens	
>	∀		ρ	∢	
LLDLTAVAGN AAVMAVIAKT PALRKFVFVF HLCLVDLLAA LTLMPLAMLS SPALFDHALF GEVACRLYLF LSVCEVSLAI LSVSAINVER YYYVVHPMRY EVRWTLGLVA SVLVGVWVKA LAMASVPVLG RVSWEEGAPS VPPHCSLQWS HSAYCQLFVV VFAVLYFILP LLLIILLVYCS MFRVARVAAM PDGPLPTWME TPRQRSESLS SRSTMVTSSG APQTTPHRTF GGGKAAVVLL AVGGQFLLCW LPYFSFHLYV ALSAQPISTG QVESVVTWIG YFCFTSNPFF YGCLNRQRG EESKQFVCFF KPAPEEELRL PSREGSIEEN FLQFLQGTGC PSESWVSRPL PSPKQEPPAV DFRIQAR	átggagtogg ggotgotgog geoggogoeg gtgagogagg teategtoot geattacaac tacacoggoa agotocgogg tgeograe cagocogogg tgeograe georgaege georgaege georgaege georgaega atchagogg cagocogoggo georgaega atchagogg cagocogogo accogogott cacacotoco atgiticogo toctgagoaga cotoacgug teggatotge tggoaggogo cagocogocotocogogo coggatotgo georgaegogo georgaegoco georgaego georgaegogo elegatogo georgaego cacacogogot elegatoga oggaaggaaggaaggaaggaaggocoggo cagocococogogo	göctcaccat ggogogoagg gggocogogo cogtotocag toggggogo acgotggoga tggoagcogo ggodggggo gtgocogo ggodggggo ggotgocogo toggogogo ggodgggo toggogogo gattgotoca ctgtottgoc ggodogogo gattgocoaggoc aaggoctacg tgctottgc cttgctcgco ttcgtggca tottggcogo gattgtgca ctctacgogo gcatcactgc caggoctacg tgctottgc ggocttgc ggocttgc ggoctgcoc ggattactg caggocogo gattactg caggocogo gattactg caggocogo gattactg caggocogo gattactg caggocogo gattactg caggocogo gattactg caggocogo gattactgc ggocttgctg cgacogotogo ggattactg ggat	IBGCCattggc caactcactt ctgaacccca tcatctacac gctcaccaac cgcgacctgc gccacgcgct cctgcgcctg gtcgctgtggg gctcccagca gtcggcggg gcggcgagg gtcccgggggggggg	FYGILANICA LYARIYCQVR ANARRIPARP GTAGTTSTRA RRKPRSLALL RTJSVALLAR VLACVLA RTJSVALLAF VACWGPLFLL LLLDVACPAR TCPVLLQADP FLGLAMANSL LNPIIYTLTN RDLRHALLR VCCGRHSCGR DPSGSQQSAS AAEASGGLRR CLPPGLDGSF SGSERSSPQR DGLDTSGSTG SPGAPTAART LVSEPAAD gttgaggeac cgtgtgtgg ccttgtocct caggccaga gcgcggcagc ccttacccc acagcgctgc agcctgcag ccttgccctcat ggcctcctat ggcctccat gg	incriagagg concoggia gogocactgo otggagggit ggraggagot otogtogoto actgggoot googgocogg ogtgaggoo agcaaggooc ggotofggt gaggaagtig gggotagaga agcagtagag cocggggooa gggacactgt tgaggiaggt gaaggocagg gagocatgga agagotgtgt goagaggtoo agggatoggo aggoggacag coagaaagoo accatgaaag coatgocaaa gateatectg gecaagaaac agaterteta gaogsocococ accateacoa
	030760		110387.1	6	
	NM_03		NP_110	LG94029	
eptor	∞		Sphingolipid Receptor Edg8	G Protein- Coupled Receptor	(HEOADS4)
Coupled GPR61	Sphingolipid Receptor Edg		Sphi	G P	图
	189900 Sphing Recept		189900 Sphi Rece	189901 GP Cou	望

Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
<u>ρ</u> .	∢	Δ.	₹ .
egecaccegg geagetgece ceaeggaage aegggteage aegtggtggg getgeaccac citeaggtag eggtgagtg egatggetgt gaggagtga etgeagetga aagagttga etgeagetg ageagecea aagegceagg gaggaagag gaggagtgg tecaegegg gggeacaggg gaggaagag gaggaagag gegaacagg gegaccagg gegaccagg agagacagg gaggacagg gaggacagg gggaccagg agatgaggag agatgaggag agatgaggag agagaccag ggccaacag ggccaggat gggaccagga agatgaagag ggccaaactg tgccaggat gggccagga ggatgagaga agatgaggg ggccaacag ggccaacag ggccaggac aaactccagg gccaggatg ggccaggag ggccagaca agatgaagag ggccaaactg tgccaggat gggccagaga agatgaagag agatgagggcc ccagaggac cccacagt ggaaaggc MELNNLSPS PSLSSVLPP SFSPSSSAP SAFTTVGGSS GGPCHPTSSS LVSAFLAPIL ALEFVLGLVG NSLAIFIECH HTRPWTSNTV FLVSLVAADF LLISNLPLRV VLSRASVGAA ARVAGGLWVG ILLNGHLLL STFSGPSCLS YRVGTKPSAS LRWHQALYLL EFFLPLALIL FAIVSIGLTI RNRGLGGQAG PQRAMRVLAM VVAVYTICFL PSIIFGMASM VAFWLSACRS LDLCTQLFHG SLAFTYLNSV LDPVLYCFSS PNFLHQSRAL LGLTRGRQGP VSDESSYQPS RQWRYREASR KAEAIGKLKV OGEVSLEKEG SSOG	ggitatggit taactcagca gaattigitg aacaactacg acatgciggg galcatggca tggaatgcaa cttgcaaaaa ctggctggtggtggtggtggtggtgggggggggg	tacattgtga cacggccttt ggcctttctg aacagtgtca tcaaccctgt cttctatttt cttttgggag atcacttcag ggacatgctg atgaatcaac tgagacacaa cttcaaatcc cttacatcct ttagcagatg ggctcatgaa ctctacttt cattcagaga aaagtgaggg gcttgtgaaa cagattgttc tacagatgaa tctgtaagcc agttacagtt tgccttaact catagacatc aatcagaga tgtcacagat ttaaccttga tctaaagaca agttgtaccc agagtatgtg aaaagaatgg gacgacaaga atgactggt tcttcctct aagaattgaa aggagttgaa ctgccttait tttgggcatg taactccaaa atactaggta gtataaggct ttctcaatca gtgcaaaaat ggaagatta, taaagcaaca agttgtctgc atttgatccc tggtcagatt gtaaaaaaaaaa	tggagccatg ctcctgggc tottccgcgg gcgcccgcgc gctgccttc gcttgaggca aaaggactct tgtggaagat ggaactcatt gtccatttc cagaatgtat ttccaagccc atcaatggga cctgatactg ctgttctgtg ttgaaatgct tgaagaactc ctgcatctct gcttgcatct tccatcctac tgaaaccatg gtcttctcgg cagtgttgac tgcgttccat accgggacat ccaacacaac
CAC38933.1	NM_033050	NP_149039.1	NM_030784
G Protein- Coupled Receptor Ls189901 (HEOAD54)	Purinergic Receptor P2U2 (GPR91)	Purinergic Receptor P2U2 (GPR91)	G Protein- Coupled Receptor GPR63 (PSP24
189901	189904	189904	189920
565	996	567	568

ggittacca aaaagctgcc atgaggtctg caattaacat ectecttgcc agectagett ttgcagacat gttgettgca gtgetgaaca

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gaaaccatg geteccactg gtttgagtte ettgaeegtg aatagtacag etgtgeecae aacaecagea geatttaaga

gocottigo colegiaaci attottacia coceategat tittegegaa ticticigia eggiatotec tatettitic tegitatite

gatagaagg agragocato otgotoatoa tragoataga taggitoott attatagtoo agaggoagga taagotaaao

ittetegtg tafgaaaaca ectacatgaa tattacacte ectocaecat tecageatee tgaeeteagt ecattgetta gatatagttt

sapiens Homo

KFFCRVSAMF FWLEVIEGVA ILLIISIDRF LIIVQRQDKL NPYRAKVLIA VSWATSFCVA PLAVGNPDL OIPSRAPQCV FGYTTNPGYQ AYVILISLIS FFIPFLVILY SFMGILNTLR

HNALRIHSYP EGICLSQASK LGLMSLQRPF QMSIDMGFKT RAFTTILILF

YYWRIKKFHD ACLDIMMPKSF KFLPQLPGHT KRRIRPSAVY VCGEHRTVV AVFIVCWAPF TTYSLVATFS KHFYYQHNFF EISTWLLWLC YLKSALNPLI

acicegaact iggeteteag egtateatec etgitaceag ggacaaatge aatticaaat titageatig giettecaag caataatgas tgottgagt catottotga agotttaaaa acaattgatg aattggoott caagatagac otaaatagca catoacatgt gaatattaca

tgagttatg tgatggcgtg cagtattgga aacattacta tccagaatct gaaggatcct gttcaaataa aaatcaaaca tacaagaact saagattotg tattagitag aagagcacag titactitot toaacaaaac tggactitic caggatgtag gaccccaaag aaaaactita cgratticc agatggatit tgagagtgga caagtggatc cactggcatc tgtaatittg cotocaaact tacttgagaa tttaagtcoa raggaagtge ateateceat etgtgeette tgggatetga acaaaaacaa aagttitgga ggaiggaaca egteaggatg gitgoacac agagaticag atgeaagiga gacagicigo cigigaacc acticacaca citiggagit cigatggacc

nalgaagict atggaaaaga aagttatggg aaagaaaaag gtgatgaatt ctgttggatt caagatocag toatatttta tgtgacotgt

acattogoc gatacattot aaaattotgo atcattggot ggggtttgoc tgocttagtg gtgtoagttg ttotagogag cagaaacaao

aguigeatt tetteettet ggeaacettt aeetggatgg ggetagaage aatteaeatg taeattgete tagttaaagt atttaaeaet

icagcoctgo tettootgaa totoototto otootagatg gotggatoao otoottoaat gregategao titgoattgo tettgoagto

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ragcagcaac tetectgaca tatgitgett ttgagaaatt gegaagggat tateceteea aaatettgat gaacetgage

itgratgigt gigagcagig taaagaaaga aiggtaatta tagitcigti accaagaata aataatagga aagigattac aaatattaco gotgatctac tactggagga ttaagaaatt ccatgatgct tgcotggaca tgatgcctaa gtcottcaag tttttgcogc agotcoctgg teatacect tectggtaat actgractea tttatgggea tacteaacae eetteggeae aatgeettga ggateeatag etaeeetgaa ccatatagag ctaaggittet gattgeagit tettgggeaa etteetittg tgiagetitt eetttageeg taggaaaooe egaeetgeag aticagiaag cacititaci atcagcacaa cittitigag attagcacci ggctacigig gcicigciac cicaagicig cattgaatco staccticce gagetececa gigigigiti gggtacacaa ecaatecagg etaecagget tatgigatti tgattietet eattiettie aaaacacgt gecticacca ctattitgat tetettiget gtetteatig tetgetggge eccaticace actiacagee tigtggcaae ccagggitc aatagaaatc ctcaaittag ggtgaggaga citititig gittiggggi tittcctiga itgattigt ittcatagig icacacaaag cgacggatac gtcctagtgc tgtctatgtg tgtggggaac atcggacggt ggtgtgaata ttggaactgg ctgacatttt gggtgatgct tgttcttat tgacattgaa ttctcttct catagcctct ccactttatt ttttttata gggtttgtgt gratatgoc teagceagge cageaaactg ggteteatga gtetgeagag acettteeag atgageattg acatgggett

gaatcagga itgigcitta itgagccigc agitacaitg aaitgiaggi gittcgigig cigciaaggi aigcitaitt gagittaica

VVCLMVYQKA AMRSAINILL ASLAFADMLL AVLNMPFALV TILTTRWIFG agactititt tittetggaa gacactgetg cittlaccat cacatiggag cc MVFSAVLTAF HTGTSNTTFV VYENTYMNIT LPPPFQHPDL SPLLRYSFET MAPTGLSSLT VNSTAVPTTP AAFKSLNLPL QITLSAIMIF ILFVSFLGNL

NP_110411.1 Coupled Receptor **GPR63 (PSP24** G Proteinbeta)

189920

569

G Protein-189945

AK027843

Coupled Receptor

Dj287g14.2

570

Seta)

gotgggtatt itggagtcat gittitictg aacaitgcca igitcatigt ggtaatggtg cagatotgtg ggaggaatgg caagagaagc

ittigeatic titigeotiggi gaccettaaa tateceette atgracetet tetecatett caatteatta caaggettat tratattea

iaccggacce tgagagaaga agtgttaagg aacetgegca gtgtggttag ettgacettt etgtigggea tgacatgggg

sapiens

sapiens

Homo

BAB55406

Coupled Recepto Dj287g14.2

G Protein-

189945

571

ttttgttoc aaggaatatg aagtgagaca tatgggtgag toataataat caaaataatt tatgaagago tgggtotgoa atagotagto icaagggaga agcaatgotg aggaagaccc tagatagagc teatittact ceaectaate gitatatetg gatataecea tittetgea soctegotoc agoagatgat gagataatga ggdagtgggt titttattac tgttocatti tgcaacatoc tgcaacaca tootgggaga acteagatig gagtaagaca getaceaata teateaagaa aagtietgat aatetaggaa aatettigte tteaagetee attggtteea acteaaceta tettacatee aaatetaaat eeagetetae eacetattte aaaaggaata gecacacaga taatgtetee tatgageatt aaaaactac tigtgigica gicctciggi tatagtatat aagagcciga ggaggicigg caagatagat ggigtattat ttatggatca nacateaate atecetetee ateaggteat tgataaggte aagggttatt geaatgetea tteagaeaae ttetataaaa atattateat getgetgea tacaaacett geatactatt atgeagetta cetaactete agactattet gagtaatget tgettgetaa tgaatgtata ragcagtgta aactgcaact agtgatgtaa atgtgctatt acctaggtaa ctgcatatat ataaggaatg tattttgtta agaaggctt attettete aacaataaae tgreettget ttggagaett taagaeattt eetaaageae aaataaaage etegtattte eeeattgaga sagaccaca tigraatigi tottagatga tggagtocat gcagtitott agaaatoggt otoagtgoat gotgigotti ticacattig gigaaatic agaatitito tititaatat atticticca iggaagagti gicatcacia aaacticagi acigagagta acatgacica sict gggtta tot gggaagt atcaggttot gggaggcaac agcattaagt gataagaaaa ggagacatto tggcaaagco cotteaacaa aagtggatea eteagacagt getteeatgg acaagteett gteaaaactg geceatgetg atggagatea natotectta aaggoaaagt coagaacote gaacotagae gootttotot otgoacgaaa aacaggtagt ttgoagtote ottecactgt getatgaagg agaatgttea gaaacagtgg eggeggeate tetgetgtgg tagatttegg ttageagata stcagacacc ttcagccaca gcacaaagtt ttaatgtctt taagaaaag aaatcaatct gcagaaatgt gaagatttgc stagecacag aagetatgat tigtaaaata tataatigaa teagagtaat cataatgeag gggagacatt caaattagag gatatggga gagcttttag gctacacagc aacceaaggg acctctcacc ttttgctgag cttcaatcag gaagctattt caagagcatt acccagcitg gcittcacgg gggagggttg tattcagt MDFESGQVDP LASVILPPNL LENLSPEDSV LVRRAQFIFF NKTGLFQDVG

gaccaaatge titgiggate tiectaccag gaatgicaae etggeecagi eegtigitat gatgaccati ggegagtiga tigggtiigi VTKVLTFISY IGCGISAIFS AATLLTYVAF EKLRRDYPSK ILMNLSTALL FLNLLFLLDG gtacatcage attgotggot ggotgateat etgeettgee tgtgtaetet tteeaeteet eagaaceagt gatgataeet etggeaalag ggitatatg aaagaaacaa aacgagctgt gatatttatg ataaacttag ccattgctga cttactacaa gttettteet tgecaetgag gatettetae taettgaate atgactggee atttgggeet ggretetgea tgitetgitt etaeetgaag tatgreaaea tgratgeaag catclactic tiggictigca tcagigiggig acgaittigg titicicatgi accoctiticg citiocatgac igcaaacaga aatatgaool agattiticga tacttiatit atgcagtgac atacactgic attottgigc caggicitcat agggaatata ttagcootgt gggrattota caccattagg caaagatagt ttetetagag agaateatge etgetaatta caegtgtace aggecagatg gagacaatae STYLTSKSKS SSTTYFKRNS HTDNVSYEHS FNKSGSLRQC FHGQVLVKTG PC KNKSFGGWNT SGCVAHRDSD ASETVCLCNH FTHFGVLMDL PRSASQLDAR /ILKFCIIGW GLPALVVSVV LASRNNNEVY GKESYGKEKG DEFCWIQDPV WITSFNVDGL CIAVAVLLHF FLLATFTWMG LEAIHMYIAL VKVFNTYIRR FYVTCAGYF GVMFFLNIAM FIVVMVQICG RNGKRSNRTL REEVLRNLRS NVOKOWRRHL CCGRFRLADN SDWSKTATNI IKKSSDNLGK SLSSSSIGSN **PORKTL VSYV MACSIGNITI QNLKDPVQIK IKHTRTQEVH HPICAFWDLN** VSLTFLLGM TWGFAFFAWG PLNIPFMYLF SIFNSLOGLF IFIFHCAMKE

572 190026 G Protein- NM_032553

Coupled Receptor

sapiens

KKRAVIFMIN LAIADLLQVL SLPLRIFYYL NHDWPFGPGL CMFCFYLKYV NMYASIYFLV CISVRRFWFL MYPFRFHDCK QKYDLYISIA GWLIICLACV

Coupled Receptor

JEG18

G Protein-

190026

Homo

sapiens

Homo

AF055084

Coupled Receptor

G Protein-

190031

574

nactocgett etgattgtee tatattgtae etggaagaeg gttttateae tgeaagataa atateeeatg geeceaagate ttggagagaa atgicitgac ceagicatat actacititic cactaatgag ticegaagae ggetticaag acaagattig catgacagea tecaacteca ggrgaagtee aatgaaatta aaagetgeet ageeagaagg gtgattetaa tattteatte tgtggeattg tgtettgeta gtergaatte egitotgoto tatottacig ctatggggaa itcacttott caaagcagga cotatitgga gcattacgat ccacgattat tgatgttgac scagaaagec tigaagatga tictaaccig igcagggga ticctaatit gettigeace trateatite agittiecti tagattiect gacctgaaat gcaagtacat cagaacatat ctgcaatacc caagccacag ggaagaactt gcaaaacaac acagcttttc gcaaaatec tttgtgagta accatacage ttecaccatg acaectgaat tatgetaaaa caaaaaacca aactgaatgt VPANYTCTRP DGDNTDFRYF IYAVTYTVIL VPGLIGNILA LWVFYGYMKE atgiccatgi agiaattitti citcaagi NP 115942.1

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igtigaagaa gaagactitig aagaacaaac tottaccott atattoctag atggagaaag agaacgtaaa gtatcagtto aaattitgga gaaccigge cagagaagea etgiatigga igicatecta aegecagaga caggatetti aaatteatti eetaaaeget teeagatigt tggccagag gcotttgtto ttoacctato aggagtgcag agoagtgoto otggoggago toaactooga toaggtttoa ttgttgotga aagactattt gggttocaca gegatettat taaagtttet tateagacea etgeaggaag egecaagoea etggaagatt ttgageetgt ggitgecat tettactgag geaactggtg tatetgecat ecctgagaaa ettgteacee tteatggeae aectgetgtg tetgaaaage ctgatgtggc cactgtaact gccaatgttt ccattcatgg aacattcagc cttgggccat ccattgttta tattgaagag gagatgaaga iggatgatac tggattigca gcttttgcca tggttattat tacagggagt gaccticaca atggcatcat aggattcagt gaggagtccc aaaagglacc acaggitgaa gigtattiti tigtggaact atatgaagst actgctggag cagcaataaa caacagtgcc agattcgcac agattaaaat cttagaaagt gatgaatctc aaagccttgt gtattittct gtgggttctc ggctggcagt ggctcacaag catectigat agitgeceat attigicaat attiggetett eactggiate eteageaaat eaatggacae aagittgaag gaaaggaagg naggicagag ticacaacte orgactaatg acaatgaggt tototacagg attiatgetg orgagectag aattatteet cagacatete teactgeag etettgitee tittgaegige etegiggigg igitegiggt giteateeat geetaeeagg igaageeaca giggaaagea natigaacca afgggegict tecaatitte cactagetea agaaatatea tagtgteaga agatacacag afgateagat tacatgtaca gtigcagig attacaatat iggataatga igacciggca ggaatggata titocitoco cgagacaact giggcigtag cagtigacac ccagittac agagtatage agccaacagt ggittataag tggaaacaat citectacce taaaaaataa ggrattatet ttgagtgtga gigicicct tiggaatcag gcigcigcaa gciggitgic igacagicag titigcaaag igatigagga aacigcagac taigiggaa gttgaggagt gctgaaacaa ttggtcgtac catcatatct ccagctattt ctggaaagga ttttgtgata actgaaggca cattggtctt paaaatteaa gettteagtg ttgecageeg aactettte tatgagatte tttgttetet tattaaccea aagegeaagg acactagggg cagaatege gaacigitti ticaaaaati ccaaactgag gtigatitig aaataaccat tattaatgat cagciticig agatagaaga rgcctgtte acacatgtet gigtatgetg tetatgeteg gaetgaeaae tigtetteat acaatgaage ettetteaet tetggattta tgazaggaa totatoatca gagoatgtoa cagatotatg gaotoattoa tggtgacotg tgtttatto caaacgtota tgotgotttg atteagteae titgetgaag tgaetgagaa tittgeetti tetetgetga etaatgitae tigeggetet eetggtgaaa aaageaaaae scial good cigicacaca traccigial citigocagi tragciggat gotcaticag icigigaati toiggracgi goiggigatg natgatgage acacagagag gegatatetg etgittitee ttetgagitg gggactacea getttigtgg tgatteteet catagitatt atgatgatg tetteagagg aaggacaaat getgeagaaa tteeaetgat tttatatete tttgetetga ttteeggae atggetttgg stittittac attaaccita citcagraga aattagggga itacaaaagi tigaigitaa tiggagcoca cgccigaatc tagatticag atgratete aggrettige tiggetgite titeceatal etietgigee aggracteea igitigeage taaactietg acteacaiga agcatgaaag tggccacaga aaacacagat gaacaactca gtgccatgat gcatctaata gaaaagataa ctactgaagg gatgatgag cotgaggggc aggaattott ctacglgttt ctcacaaacc ctcaaggggg agcacagatt gtggagggg igattacatt cgaattecag agaggetact ggatgtecag gatgeagaaa taatggetgg gaaaagtaca tgtaaattag gaagatgica aggictitig gogagicaca citaacaaaa cagicgicgi golocagaag gaiggggiaa accigaigga a actoto att cot graga a a cega atoca cacatacoto ago a caga ga o ga o ga cotto tego ago co a aco a co igagiggact agaactcagg gaaggagctg ttatgagaag attgcacctt attgtcacaa gacagccaaa cagggcctt ggaacticag tetgigteag ggaceaeaae etgiacaatg ggicaaaeaa aatgettiat eageattgaa eteaaaeeag cgcaggccat ttgggggctt gcagatcagc tacatcagcc tgtgaatgat gatattctca acagagtgct ccataccatc ggcagccag citaggiaca cagaitctgt ttctggcgtc tgcatacgca agtccccaac tcgctgagga gagctgttca aaggecaetti taateagtet geaggtggee agagattetg ggacaggaet aatgatgtet gttaaettta gtaeceagga itggcacatt caacactgca gaagttetta teegaagaac tggtgggttt actggcaatg teagcataac agttaaaact teggtgaaa gatgtgetea gatggaacea aatgeattge cetttegtgg tatetatggg atttecaace taacatggge ccittitigac ccaaaaggig gigccagaat igataaagig taigggacig ccaacatcac tctigictca gaigcagait

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DLITNDNEVL YRIYAAEPRI IPQTSLCLLW NQAAASWLSD SQFCKVIEE

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JSDADSQAIW GLADQLHQPV NDDILNRVLH TISMKVATEN TDEOLSAMMH

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geacacttt catattigta teagetttig igetaaaact etetaagtae atecacetgt gtaataggaa eelgtgaatt gtaetggatg

sactgactee cagateging ageteaggag gatacecate geogacaete accigtagea ceteactaae cattegactg tattigcat taaaaacigg tgctggtctc agtgtcagtg ataatgaatc tggtcaaggc agccaggagg ggggcacctt

AAD55586.1

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VFLSLGSNFT LOLVTVMLVG GRFYGMPTIL OEAKSAVLPV SEKAANSOVG

FESTAFOLMN ITAGTSHVMI SRRGTYGALS VAWTTGYAPG LEIPEFIVVG

LRSGFIVAEI EPMGVFOFST SSRNIIVSED TOMIRLHVQR LFGFHSDLIK VSYQTTAGSA KPLEDFEPVQ NGELFFQKFQ TEVDFEITII NDQLSEIEEF FYINLTSVEI RGLQKFDVNW SPRLNLDFSV AVITILDNDD LAGMDISFPE TTVAVAVDTT LIPVETESTT YLSTSKTTTI NMTPTLGSLS FSHGEORKGV FLWTFPSPGW PEAFVLHLSG VOSSAPGGAO COPTINIVATIV TEATGVSAIP EKLVILHGTP AVSEKPDVAT VTANVSIHGT FSLGPSIVYI EEEMKNGTFN TAEVLIRRTG GFTGNVSITV KTFGERCAOM KDGVNLMEE LOSVSGTTTC TMGQTKCFIS IELKPEKVPO VEVYFFVEL DEPEGOEFFY VFLTNPQGGA QIVEGKDDTG FAAFAMVIIT GSDLHNGIIG SEESOSGLE LREGAVMRRL HLIVTROPNR AFEDVKVFWR VTLNKTVV EPNALPFRGI YGISNLTWAV EEEDFEEQTL TLIFLDGERE RKVSVQILDD ZATAGAAINN SARFAQIKIL ESDESQSLVY FSVGSRLAVA HKKATLISLQ PGORSTVLDV ILTPETGSLN SFPKRFOIVL FDPKGGARID KVYGTANITL VARDSGTGLM MSVNFSTOEL RSAETIGRTI ISPAISGKDF VITEGTLVFE

Coupled Receptor G Protein-**VLGR1** 190031

	Homo	Homo sapiens	Homo sapiens	Homo sapiens
·	∢ .		∢	Д.
ADYVECACSH MSVYAVYART DNLSSYNEAF FTSGFICISG LCLAVLSHIF CARYSMFAAK LLTHMMAASL GTQILFLASA YASPQLAEES CSAMAAVTHY LYLCQFSWML IQSVNFWYYL VMNDEHTERR YLLFFLLSWG LPAFVVILLI VILKGIYHQS MSQIYGLIHG DLCFIPNVYA ALFTAALVPL TCLVVVFVVF HAYQVKPQW KAYDDVFRGR TNAAEIPLL YLFALISVTW LWGGLHMAYR HFWMLVLFVI FNSLQGLYVF MVYFILHNQM CCPMKASYTV EMNGHPGPST AFFTFGSGMP PAGGEISKST QNLIGAMEEV PPDWERASFQ QGSQASPDLK PSPQNGATFP SSGGYGQGSL IADEESQEFD DLIFALKTGA GLSVSDNESG QGSQEGTLT DSQIVELRRI PIADTHL	aggratical traiggoagg atocatatic arcacaatat tiggoaatot tgocatgata attitocatit octaciticaa goagotticac acaccaacca actitoctoat cototocaig gocalcactig attitoctoct gggaticaco atoatgocat atagtatgat cagatoggg gagaactaco atoatgocat atagtatgat cagatoggg gagaactacot gagaactacot gagatocatit tecagiggoca tigatagatit tatgotaat tatagatit atatagiti teacotgatig citagocata calcattiti tocattigo toagggoca tigatagatit tatgotata tetitatic caccaaata aclaticoag toattaaaag attgotactit calgitigg citagitigo togggocat tacitatatic caccaaata aclaticoag toattaaaag attgotactit cactotggg gicatgatig gegggocate togggagacac citatitatig geaggitict toactotgg gicatgatg gegggagatag gattictig geaggitict toactotgg gicatgatg gegggagatag gattictit attagtigg titoctigit totacaataa titatagga accitatiga actitotoac tocigaagit tigtigatig octigacag gagtiticti attagtigg titoctigit totacaaat titatiggat coctititiga actitotoac tocigagit tigtitgatig octigacagi gagattictit attagtigg titoctigit tocataatac gitaatatat agittictot alocotggit togcagagaa otgaagaaca tittgoragg taaaattito agotoatgit tocataatac tattiteria atecaaaaa aaaataagaa	MYSEMAGSIF ITIFGNIAM ISISYFKQLH TPTNFLILSM AITDFLLGFT IMPYSMIRSV ENCWYFGLTF CKLYYSFDLM LSITSIFHLC SVAIDRFYAI CYPLLYSTKI TIPVIKRLLL LCWSVPGAFA FGAVFSEAYA DGIEGYDILV ACSSSCPVMF NKLWGTTLFM AGFFTPGSMM VGIYGKIFAV SRKHAHAINN LRENQNNQVK KDKKAAKTLG IVIGVFLLCW FPCFFIILLD PFLNFSTPVV LFDALTWFGY FNSTCNPLIY GFFYPWFRRA LKYILLGKIF SSCFHNTILC MQKESE	atggatctaa citalatice cgaagaceta iccagitgic caaaattigt aaataagate cigtocicce accaaceget citticatgi ccaggigata atgaticeg italgactgg agecatgati alceaciati cggaaactig gitataatgg iticcatate geaticaaa cagcitcaci ciccacaaa citticigate ciclocatgg caaccacgga citticigcig ggittigica tatigicata cagcataatg cgatcagtgg agagtgetg gategottit glaaattoca cacaagctti gacatgatge teagactgac ciccattic caccitcigt ceatigciat tgacegatti tagacegitti glaaattoca cacaagctti gacatgatge teagactgac ciccattic cactegit catigcigat atgacegatt tagacegitti gracetti acataacaa accaaaatga cgaactccae cataaagcaa ctgctggcat titigctgac citacttica acaaattcig ggggacaata tititicacta catgiticit taccctggc atactigtig citicataa titiciggce citacttica acaaaattcig ggggacaata tititicacta catgiticit taccctggc teagagaaaa acacciatic caagaaaaate tatacggac acgaagaaa acteagagaa acagaagaga cagegaagaa acteagggata gtategggg tititicggc titiciggtig cettigtitic tagatgaccca tacctagact accaactac catacaata tiggatcitt tagtggggt cottgataaa cacaaaagaaa accaactcact gaaaaccctt tatactage titicagaca acteaacaa catacaata tiggatcitt tagtggggataaaaaaaccaactit caagaaaccctit tagtatagge titicagaaa accaactaca tigaacctat gaaaacccto catacaaaa catggittica gaaagacattc aagacaaaaa gacaagaaaaaaaaaa	addings, tocalicae agartigcae inglitor gaggacatil ag MDLTYIPEDL SSCPKFVNKI LSSHQPLFSC PGDNVFGYDW SHDYPLFGNL VIMVSISHFK QLHSPTNFLI LSMATTDFLL GFVIMPYSIM RSVESCWYFG
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	G Protein- Coupled Receptor GPR58	G Protein- Coupled Receptor GPR58	G Protein- Coupled Receptor GPR57	G Protein- Coupled Receptor
	190168	190168	190170	190170
	<i>S</i> 76	577	578	. 625

gaaccacttt gggaaccccc aaccctccat ggatggagaa ctgctgctga gggcagaggg atctacgcca gcaggtggag

sapiens

secactgeca ggaggaegge ateatgetgt etgeegaetg etetgagete gggetgteeg eegtteeggg ggaeetggae

AB049405

Coupled Receptor G Protein-

190188

LAFCWSVPA LFSFGLVLSE ADVSGMQSYK ILVACFNFCA LTFNKFWGT

DGFCKFHTSF DMALRLTSIF HLCSIAIDRF YAVCYPLHYT TKMTNSTIKQ FITCFFIPG SIMVGIYGKI FIVSKQHARV ISHVPENTKG AVKKHLSKKK

ORKAAKTLGI VMGVFLACWL PCFLAVLIDP YLDYSTPILI LDLLVWLRYF

NSTCNPLIHG FFNPWFQKAF KYTVSGKIFS SHSETANLFP EAH

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cataatcate tggettitet teetgeagtg etgeateeae ecetatgtet atggetaeat geacaagaee attaagaagg aaateeagga gtaacagca acagcaaccc tectetgecc aggtgetaec agtgeaaage tgetaaagtg atetteatea teattitete etaigtgeta ggitagcete acceacetgt tegeettege cagegteaae aceattgleg tggtgleagt ggategetae ttgteeatea tecaecetet <u>මුපුපුපෙහුලෙ</u> අපුළාඅපූරයෙනු අයුතුළතුදෙන් අරමුදැල්පුරෙන පුරසුතෙල්ලියනු පෝඩුළත්ළුපූව තතුළත්තමුදය සුප්තරයක්ක්මු gggcagaat ggaagccaag gacggcagcc tgaaggccaa ggaaggaagc acggggacca gtgagagtag tgtagaggcc igetacacta ttotcagogt ggtgtoctto atogtcatto cactgattgt catgattgcc tgctactoog tggtgttotg tgcagocogg igagggagca gagaagaagg aggagttcca ggatgagagt gagtttcgcc gccagcatga aggtgaggtc aaggccaagg gaggagaac agcatgaagg cagacaaggg tegeacagag gteaaceagt geageattga ettgggtgaa gatgacatgg atgacgteca cotgeaceaa cageaegege gagagtaaca geagecacae gtgeatgece etetecaaaa tgeceateag ratgotgaag aagttottot goaaggaaaa gococogaaa gaagatagoo accoagacot goooggaaca gagggtggg iggcagcaig cicigcigia caaigicaag agacacagci iggaagigcg agicaaggac igigiggaga aigaggaiga igittigitga agacgacato aatticagtig aggatgacgt cgaggcagtig aacatcccgg agagcotoco acceagtogl ctectacceg tecaagatga cecagegeeg eggttaectg etectetatg geacetggat tgtggeeate etgeagagea stoctocact ctaeggetgg ggecaggetg cetttgatga gegeaatget etetgeteca tgatetgggg ggecageeee ccotgggge cotactgott tttagcagte ctggccgtgt gggtggatgt cgaaacccag gtaccccagt gggtgateac cotggoccac ggcatcatcc gotcaaccgt gotggttatc ttoctogoog cototttegt oggcaacata gtgotggogo tagitgitgca gogoaagcog cagotgotgo aggitgacoaa cogititato titaacotoc fogicacoga cotgotgoag attrogeteg tegecocote getegetege acctetete cietettote gecoteaac agocactict geacegeect otgaaggcaa gattgtccct tcctacgatt ctgctacttt tccttga

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Coupled Receptor G Protein-

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Homo sapiens

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G Protein-Coupled Receptor

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RRVKGFHSCC YDCVDCEAGS YRQNPDDIAC TFCGODEWSP ERSTRCFRRR

SRFLAWGEPA VLLLLLLSL ALGLVLAALG LFVHHRDSPL VQASGGPLAC

FGLVCLGLVC LSVLLFPGOP SPARCLAOOP LSHLPLTGCL STLFLQAAEI

*VESELPLSW ADRUSGCLRG PWAWLVVLLA MLVEVALCTW YLVAFFPEVV

IDWHMLPTEA LVHCRTRSWV SFGLAHATNA TLAFLCFLGT FLVRSQPGRV

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LRSVLSSFA AALCEERPGS FIPTEPQTQL DSEGPTLPEP MAEAQSOMDP

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WVLATLFSV PWLVFPEAAV WWYDLVICLD FWDSEELSLR MLEVLGGFLP ŁYYFLWGVS YSSGLFLLAA LSLDRCLLAL CPHWYPGHRP VRLPLWVCAC JAOLLYLAFL WDVYSGYLLW EALVYSDYLI LLNSCLSPFL CLMASADLRT TLLLVCHVL TQATACRTCH RQQQPAACRG FARVARTILS AYVVLRLPY ENSMPRT2619 Coupled Receptor G Protein-.s190484 190484

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Homo sapiens

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Homo

599 190599 G Protein- NP_057319.1 Coupled Receptor GPRC5B

190602 G Protein- NM_014373 Coupled Receptor GPCR150

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Homo sapiens

sapiens Homo acccagecat ctaccaaage ctgaaggeac agaatgetta ttetegteac tgteetttet atgteageat teagagttae tggetgteat ctttggatcc atttgtcaac tggaagtgct getteattec acttacaatt ectaatettg agcaaattga aaageetata teaataatga aaaaacaaaa taattocaag aagttittat agttattoag ggacaciata ttacaaatat tacttigtta ttaacacaaa aagtgataag agitaacatt tggctatact gatgtttgtg ttactcaaaa aaactactgg atgcaaactg ttatgtaaat ctgagatttc actgacaact ttigitaata itattaatta aaagttacag cigicataag atcataatti tatgaacaga aagaactcag gacatattaa aaaataaact RITSYMNET ILYFPFSSHS SYTVRSKKIF LSKLIVCFLS TWLPFVLLQV IIVLLKVQP gaactaaaac aacttttgcc ccctgactga tagcatttca gaatgtgtct tttgaagggc tataccagtt attaaatagt gttttatttt CONFINEYFCI SLAFVDLLLL VNISIIL YFR DFVLLSIRFT KYHICLFTOI ISFTYGFLHY tticatggt gatgattita titgtagctt tcataaccig ttgggaagaa gitactactt tggtacaggc tatcaggata acitectata gitticicag tacciggita ccattigiac tacticaggi aatcatigit ttactiaaag ticagaticc agcatatatt gagatgaata gaatgaaac tatettatat titeetitti eateeceacte eagitataet gigagateta aaaaaatatt ettateeaag eteatigtet tecctegit atactitigte aatagitite teatigetae agigiatigg ittaatigte acaagettaa titaaaagae atiggatiae YOSLKAQNAY SRHCPFYVSI QSYWLSFFMV MILFVAFITC WEEVTTLVQA AYIEMNIPWL YFVNSFLIAT VYWFNCHKLN LKDIGLPLDP FVNWKCCFIP MTALSSENCS-FOYQLRQTNQ PLDVNYLLFL IILGKILLNI LTLGMRRKNT PVFLTACIDY CLNFSKTTKL SFKCQKLFYF FTVILIWISV LAYVLGDPAI taagatate aacetaaaca tttttattaa atgtteaaat gtaageaaga aaaaaaaaa

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Coupled Receptor

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sapiens		Homo			Homo	Homo sapiens
GTWAAAWVPL PTVDVPDHAH YTLGTVILLV GLTGMLGNLT VIYTFCRSRS LRTPANMFII NLAVSDFLMS FTQAPVFFTS SLYKQWLFGE TGCEFYAFCG ALFGISSMIT LTAALDRYL VITRPLATFG VASKRRAAFV LLGVWLYALA WSLPPFFGWS AYVPEGLLTS CSWDYMSFTP AYRAYTMLLC CFVFFLPLLI IIYCYJFFR	PYSAVALVAF AGYAHVLTPY MISSVPAVIAK ASAIHNPIIY AITHPKYRVA IAQHLPCLGV LLGVSRRHSR PYPSYRSTHR STLTSHTSNL SWISIRRRQE SLGSESEVGW THMEAAAVWG AAQQANGRSL YGQGLEDLEA KAPPRPQGHE AETPGKTKGL IPSQDPRM	atggalacag geocogacca groctaette teeggeaate actggitegt etteteggig taecitetea ettroetgig ggggeteece. A cleaacetge tggeodgig ggettegig gggaagetge agegeogoc ggtggoegig gaeggerice tgeteaact gaegocoteg gaeggetee tgetgetgit etgecttie egeatgigg aggaagecaa tggaatgae tggeocoteg ettlediet etgettiene energian ethertaet ethere et etgeteet etgeteet ettlediet energian ethere et	loctgagtgt egoccacca cigigetaca agaccegac gaggetgggg caggcaggte tegtgagtgt ggoctgetgg cigitagect cigitagect cigitagect cigitagect gagetggt tagtcaggg cagcaggte tagtgagtg ggoctgetgg gaccigetgg gaccigetgg gaccigetgg gaccigetgg gaccigetac gaaggacaca getagecate cicitagecat taggetgga gaccigetgg gaccigetgg	legicocycu galcalicator agotatorgot acagoogoot ggrgrggalo cioggoagag ggggcagoo cogooggoag aggagggggg oggggotgit ggoggocaog otgotcaact toottgictg cittggggoot tacaacgrgt cocatgrogt gggotalato tgoggtgaaa gcooggoatg gaggatotac grgaogotto toagcaooot gaactootgt grogaoood tigtotacta citotoctoc toogggitoc aagcogacti toatgagotg otgaggaggi tigtgigggot cigggggocag tggagoagg agagoaggat ggagotgaag gagoagagg gagoggaggagga gaagagagg chaogaaga	gaccagigaa cactcacagg gcigtggaac tggtggcaa gtggccigg ctgaaagcia g. MDTGPDQSYF SGNHWFVFSV YLLTFLVGLP LNLLALVVFV GKLQRRPVAV DVLLLNLTAS DLLLLLFLPF RMVEAANGMH WPLPFILCPL SGFIFFTTIY LTALFLAAVS IERFLSVAHP LWYKTRPRLG QAGLVSVACW LLASAHCSVV YVIEFSGDIS HSOGTNGTCY I FFRR TDG 141 11 PVR 1 FMA V VII FYAD 1 11 11 PVR 1 FMA V VII FYAD 1 11 11 PVR 1 FMA V VII FYAD 1 11 11 PVR 1 FMA V VII FYAD 1 1 11 PVR 1 FMA V VII FYAD 1 1 1 PVR	SYCYSRLVWI LGRGGSHRRQ RRVAGILLAT LLNFLVCFG VIVER CONTROLL OF CONTROLLAND CGESPAWRIY VILLSTLNSC VDPFVYYFSS SGFQADFHEL LRRLCGLWGQ WQQESSMELK FQKGGEFQRA DRPAERKTSE HSQGCGTGGQ VACAES caagactgct cototogcc gactacaca gattggagc atggcttigg agcatagc agcatagact acagtcaata tgactgac tgatcaaag aagatacag aatatatag targant acagtcaata tgactgac tgatcaaag aagattcag aagatttcc
		G Protein- NM_005304 Coupled Receptor GPR41 & GPR42			G Protein- NP_005295.1 Coupled Receptor GPR41 & GPR42	C-C Chemokine NM_016557 Receptor 11
		604 190627			605 190627	606 190701

receptoric cototogo gactacaca gatigago algottiga agragaaca graacagat tattattatg
aggaaaatga aatgaatgo acttatgact acagtcaata tgaactgat tgatcaaag aagaattgoa aaagtttoc
tecotgatt cotoacaata gittogoa acttatgact aggcaattoc atgataggg caattatgo cattacaag aaacagaga
coaaaacaga tgtgacato etgaattgg otgagcaga ttactoctt cattcacto tgoctttta ggctgtaat gcagttoatg
ggtgggttt agggaaata atgtgacaaa taacttcago etgacaca ctaaacttg tototgaat gcagttotg gettgatoa
gcatagacag atatgtggca gaactaaag tococagoca alcaggagt ggaaaaccat gctggatcat ctgtttctgt
gtotggatgg otgcatott gctgagcata coccagctgg tittitatac agtaaatgac aatgctagg gcattoccat titococgc
taccaggaa catcaatgaa agcattaaa agatotgcat tggatttga gaccottto ttattatggg ggtgctac
titatcacag caaggacact catgaagatg coaaacatta aaatatotog accoctaaaaa gttctgctca cagtgctata agtittcatt
gfcacaac tgcottataa cattgccaag ccatagacat catctactoc otgatcacca gctgcaacat
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gagcaaacgc atggacatog ccatccaagt catcgcactot ttcacagctg cotcaaccca atcottatg

sapiens

/FVIGLAGNS MVVAIYAYYK KQRTKTDVYI LNLAVADLLL LFTLPFWAVN

MALEQNOSTD YYYEENEMING TYDYSQYELI CIKEDVREFA KVFLPVFLTI

NP_057641.1

C-C Chemokine Receptor 11

607

NM_016568

Coupled Receptor

SALPR

G Protein-

190705

809

Homo

sapiens

Homo

GKPCWIICFC VWMAAILLSI PQLVFYTVND NARCIPIFPR YLGTSMKALI QMLEICIGFV VPFLIMGVCY FITARTLMKM PNIKISRPLK VLLTVVIVFI VTQLPYNIVK FČRAIDIIYS LITSCNMSKR MDIAIQVTES IALFHSCLNP ILYVFMGASF KNYVMKVAKK icgregetge agetteegga ettgtggtgg gagetgggge tggagttgee ggaeggegeg eegeeaggae ateeeeggg Iggcagcagg cggggacaag ctagcagaac tcttcagtct ggtcccggac cttctggagg cggccaacac gagtggtaac -agoggoggg goagagagog oggacacaga ggooogggtg oggattotoa toagogtggt gtaotgggtg gtgtgogoo gatttgggga gttatgcgcc agtgccccag tgaccgcggg acacggagag gggaagtctg cgttgtacat aaggacctag ggragcogga ggacgcccga ccggagccag cgcccggaga ctgtcgaagg tcaccaaatc agtgaccatc gttgtcctgl teccatgag tetgacgege taccattegg tegectegge tetgaagage caceggacce gaggacaegg ceggggegg gecateatta tettigtigeta eetgetgetg gtgegettea tegeegaceg eegegeggeg gggaceaaag gaggggeege egictigic ecceagaaca igacetagag graceigoge atgeagatgg eegatgeage caegatagee accatgaata recegitgag cteaactect gegtecaggg egitegetge gegecaggae gegetiagia eccagitient gggetetete ggootogotg cocagtgoca tittotocac caeggicaag gtgatgggog aggagotgtg cotggtgogt ttocoggaca igitigatigig cagagacagg cagitatigiga tiggigactota acactogaag aaggtigatigi tiggigatiogi gatgoogatig gactocgag cttggcctga gaaccettgg acgocgagtg cttgccttac gggctgcact cctcaactet getecaagc gctgcggcc ggagcctggg ggacagctgc tgcttctcgg ccaaggcgct gtgtgtgtgg atctgggctt tggccgcgc ectetactge etegtgégee gegagtteeg caaggegete aagageetge tgtggegeat egegteteet tegateacea gggttggc gggcaacctg ctggttctct acctgatgaa gagcatgcag ggctggcgca agtcctctat caacctcttc caggagiatt teetgigeea ggtataegeg tteeetgiga gegigigeet agegeaetee aaeagetgee teaaeeegt icagiagci gottigaaag cicccacgca cgicccgcag gciagccigg caacaaaaci ggggiaaacc gigitaicti cettetteet grgtiggetg eccaaccagg egeteaceae etggageate eteateaagt teaaegeggt geoetteage iggocotto ggeaaggeea igigiaagai ogigiocaig gigaogicea igaacaigia egecagegig itettocica caccaacc tggcgctgac ggactticag tttgtgctca ccctgccctt ctgggcggtg gagaacgctc ttgacttcaa AVHGWVLGKI MCKITSALYT LNFVSGMQFL ACISIDRYVA VTKVPSQSGV YGSWRRQRQS VEEFPFDSEG PTEPTSTFSI

sapiens

Ното

sapiens

Ношо

aaaaagcatg cagaaaaaga agcagacgtt ttacattggg aattaatgaa agcgtgtctg ctagttttgg gtaggagaac

Coupled Receptor GPR85 (SREB2)

ADAA ITA TMINKAAGGDK LAEI ELPDGA PPGHPPGSGG AESAD LAKKSMQ GWRKSSINLF VTNL, SLGDSC CFSAKALCVW IWALA LLGRDR QFWLGLYHSQ KVLL, GAAVAG GRPTGASARR LSKV JAVPFS QEYFLCQVYA FPVSVC WRIASP SITSMRPFTA TTKPEHI YSGGRY DLLPSSSAY	CGGRSLGRDR OF THE LANGE AND LLEAANTSGN ASLOLPDLWW ELGLELPDGA PFGHPGSGG AESADTEARV RLISVYWV VCALGLAGNL LVLYLMKSMQ GWRKSSINLF VTNLALTDFQ FVLTLPFWAV ENALDFKWPF GKAMCKIVSM VTSMNMYASV FFLTAMSVTR YHSVASALKS HRTRGHGRGD CCGRSLGDSC CFSAKALCVW IWALAALASL PSAFSTTVK VMGEELCLVR FPDKLLGRDR QFWLGLYHSQ KVLLGFVLPL GIILCYLLL VRFIADRRAA GTKGGAAVAG GRPTGASARR LSKVTKSVTI-VVLSFFLCWL PNQALTTWSI LIKFNAVPFS QEYFLCQVYA FPVSVCLAHS NSCLNPVLYC LVRREFRKAL KSLLWRIASP SITSMRPFTA TTKPEHEDQG LQAPAPPHAA AEPDLLYYPP GVVVYSGGRY DLLPSSSAY GGVVVYSGGRY DLLCAGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG	NM_018970	G Protein- NP_05/652.1
とりしゅく ロンドック st	talis daliji se se se se tito at listere pe se alije ateks se se. Sije tito	NP_057652.1	NP_057652.1 seceptor NM_018970

609

ttgggtitica taalaggagt cagoglggtg ggcaaoctoc tgatotocat titggcagtg aaagataaga cottgcatag agcaocitac tacttoctgt tggatotttg otgttoagat atootcagat otgcaattig titoccatit gtgttoaact otgtoaaaaa tggototaco actetigicity tegecatege attreccee gittiagaes teggeacita eteaticati agggaggaag ateaatgeae ettecaacae ccagggggat tictaacage tgctgfctgg;atgagttig cccaagcagg aatcaatcci titgtctgca titictcaaa cagggagctg gaaagtagc aggtgctaag tatcagtgct aaatgctctg tatgtcacta catatgaaaa aacatcaaaa aacaattagc attggacatc ctocatocat ctatégogaa ctatagocat goagotgaca acattitgoa aaatototog cototaacag cottiotgaa aotgaottoo gicaccagat acitagotat egeceateae egettetata caaagagget gacettitigg aegigietigg etgigatetg tatggigigg iggaagtig itgettaaaa ittiatatea eeteeacaaa eaaaaetett eggaaatggt aaaataagaa aatgeatgat tetagagge catetgiaaa tettiageet tetgaaaaact aacettetet getgageaat tetggeeeat agecatatti tgagaagaaa tteaagaatg tttgcaaaga ctaaaatatt tggggactta aagtactgta atccactaaa gacgtgccaa tgaattattg gaatatcaca ctttaaaaac ttoctaagoa cocaceteto aggottigig gigioteteg atacatooga coettiggao iggitagggo tracigagag otocattio egetectica gggetaatga tteettagga titatgetge tiettgetet eatecteeta gecacacage tigtetaeet eaagetgata ictatataa igacittict gittetaace itgiggggee ectaectggi ggeetgitai iggagagitti itgeaagagg geetgragia gaatcagcag ttttaaggat ttgggcaaca ttctgcagtc tttgcaatag ttcacctata atcctatttt aaatctcaga gtgatcctgc Egacitate geactoteac tigicazagie attectitic iggegettit giccigittic cacactecti toatectot ofecatore taataaatt aagttgacat gaggtaaatg tgttgataaa aactaatttt agaagtttga agactttaaa acatttcata ctactattgt egectigiaa giteiggga geaticeaaa geagatati ggiteeaati agagtitaet tittigat taataeatig etattietaa iggaitatat titicagraaa atgratggat ctatctitic ctigiticita tatciagatc atgagactig actgaggctg tatcctiatc ggoaaaatgo aaacaccaca ggoagaagaa ggotattggt ottagaogag ttoaaaatgg agaaaagaat cagoagaatg egagocagt ggocaggoag otgocaattg gotagoagga titggaaggg gtocoacaco accoacttg otgggoatca ggaaagoott acaagactga ggaatatcag actgcgaatc accgggaacg gttcottigc agcacagaag caatctctct aggogotgtt tcagcacaac cottetttae tgcagaaaat ccaggttace aagggaacet tactgtgtta tatgagggag titticgice aegategaag aaaaatgaag eeagteeagt tigrageage agteageeag aactggaett tieatggiee gactgecag caaaggtttg taattaagaa gggactgaac cactgeceta agtttettta tgtggteaaa aactagataa

	Homo sapiens	Homo sapiens Homo sapiens	Homo sapiens
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ataccactit cotoatotac tagtaagatt gotagcattg aactgatta tytggtittit gitgattigg tataaagtit ticcaattoa titatattit acaaatgota gatattggt taggaggaa cattaattggt accagcotgt cacaactgag cagtictaat aatgagaat aaalacatgt tectaaag ggtatotag tatottoat citatitage actgaggaa atagccaagg gaaatcaaat cagtaactgg categatat gatataaa ggtatotag tatottoat citatitage actgagagaa atagccaagg gaaatcaaat cagtaactgg cagtotaaa ggtaatotaaa gatcattat tactitic tittitict acatggittg aaacttaaag tgcacacac (gaaataagg gaagtcaaag gaattaaaa gatcattat tactitic taggaggag gaggittat attitaagtca gctgtcaagg gaagtcaaag gacactaa gctgtcaagg gaagtcaaggag gaagtcaaggag gaagtcaaggag gaagtcaaggag gaagtcaaggag gaagtcaaggag gaagtcaatt caggtatit agaattaaa tatiticagt titaactigt gaagctta attatgatt citatgatti agaaatacat totggattig gagtctatt citaagata cagattgrg aacticaata taaagttggag gcaagccaaa tataccgg tagcctgtta attitictga aataagttig cittitigga cataacaa cettitititit aatitggag gcaagccaaa actaggaaga cagcttat aattatictg taaaagcatt attataaat calcocicta tiatictaa atgcaagaa tataagga aactgaata aacatcaacac titaattigg agcatagaac catagaaatt tiggggticta aatatacaac titaataata aacatcaacac titaattigg agcatagaac catagaaatt tigggticta aatatacaac titaattigg agcatagaac acagaaatt tigtitigt tiatiggitig gittitigga agtitatiga agtitatit tiatitggitg gittitigga agtitatiti tittitgga gaataaaaa aaaaaaaaaa	MÄNYSHAADN ILONI, SPLTA FLKLTSLGFI IGVSVVGNLL ISILL VKDKT EHRAPYYFLL DLCCSDILRS AICFPFVFNS VKNGSTWTYG TLTCKVIAFL GVL.SCFHTAF MLFCISVTRY LAIAHHRFYT KRLTFWTCLA VICMVWTLSV AMAFPPVLDV GTYSFREED QCTFQHRSFR. ANDSLGFMLL LALILLATQL VYLKLIFFVH DRRKMKPVQF VAAVSQNWTF HGPGASGQAA ANWLAGFGRG PTPPTLIGIR QNANTTGRRR ILVLDEFKME KRISRMFYIM TFLFLTLWGP YLVACYWRVF. ARGPVVPGGF LTAAVWMSFA QAGINPFVCI FSNRELRRCF STTLLYCRKS RLPREPYCVI	aggiciaging agricitic cardinator artigogoco artigogono aggiciagina agriciagina acagicagoco artigogoco artigococo artigococo artigococo acidococo acidococo acagicago acidococo acidococido acidococo acidococo acidococo acidococo acidococo acidococo acidococo acidococo acidococo	atggccaaca ctaccggaga gcotgaggag gtgagcggog ctotgtoccc acogtcogca tcagottatg tgaagctggt actgctggga ctgattatgt gcgtgagcct ggcgggtaac gccatcttgt ccctgctggt gctcaaggag cgtgccctgc
	NP_061843.1	LG93120 LR26	NM_018969
	G Protein-Coupled Receptor GPR85 (SREB2)	G Protein- Coupled Receptor GPR26 G Protein- Coupled Receptor GPR26	Sreb3
	190711	190725	190741
	611	613	614

SCKEINRIL HRRSIHSSGI TODSHSQNIL PVSE
atggccaca cregicagaga-gcctgaggag gtgagoggog cictigicoc acogicogca tragettaig tgaagctggt
acigctggga cigattaig gogtgagcct gggggaac gcatctigt ccctgctgg gctcaaggag cggccctgc
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tattcgggag gaggaccaff gcatcttiga gcatcgctac ttcaaggcca atgacacct gggcttcatt
tattcgggag gaggaccaff gcatcttiga gcatcgctac ttcaaggcca atgacacct gggcttcatt

Unidentifi

ggcagctacc catgctgtot acggcaagct gctcctctte gagtategte accgcaagat gaagccagtg cagatggte cagatgtes cagatgtes cagatactg groceggge caccggcag gctgcaca actggatege eggctttggc cagacateg groceggge caccggcag gctgcaca actggatege eggctategg gcatggacga gtgtggcca tgccaccaac eggctactgg gcatggacga ggtcaagggt gaaaagcact gccaccateg gttcaagggt gaaaagcgt gtgtgacgt gacacaggc tetttcigct ectetggtea ecctacateg ggtcaagggt gttgtgaaag ctgtgcgt gcccaccgc tetttcigct ectetggta ecctacateg gggcgta ttgtgaaag ctgtgcgt gcccaccgc tetttcigct ectetggta cctacateg gggcgta eggcgacta ggaggttc tettgcta acaaggact caagaagtgc etgaggact acgggcaca ggaggtgcc eggctccag agaacctac tgtgtcatg ga MANTIGEPEE VSGALSPPSA SAYVKLVLG LIMCVSLAGN AILSLLVLKE PAALHKAPYYF LLDLCLADGI RSAVCFFVL ASVRHGSSWT FSALSCKIVA FMANTICEPEE VSGALSPPSA SAYVKLVLG LIMCVSLAGN AANICAAT HAVYGKLLLF EYRHRKMKPV QMVPAISQNW TFHGPGATGQ AAANWIAGFG RGPMPPTLLG IRQNGHAASR RLLGMDEVKG EKQLGRMFYA ITLLFLLWS PYIVACYWRV FVKACAVPHR YLATAVWMSF AQAAVNPIVC FLLNKDLKKC LRTHAPCWGT GGAPAPREPY CVM

NP_061842.1

Sreb3

615

E32367

G Protein-Coupled Receptor

H7TBA62

Homo sapiens

> eletelecet ggggtggagg ettggggetg ecotecatag eggggtaaet etecettete ecotecetet etgecattta gageoeetet aagcactiaa ttotacagcc tecttectag agcetteagt ggeetetgee agtetggeag acactigeag acetetette teagcacea scaaccatgt ggtcactoto tggggtgtoc tggtgaagtt tgacotggtg ocotggaaca gtactttota tactatocag aogtatgtot cceteggeat ggtaatagee teteagtace ettetgeeae aaacaceeea aactteteet ttgaaataat atteatacaa attgetatt aggaacagaa ctgagggcat gcccaggtcc acacaggccc tcataggccc agtgttccca gtggggagga aacaggaagc igigacitics tetetetiti ecetecetge tettageete aaggteactg etgetgagat gaatteeaae etgittiagt tggeaetgit aggototgg caggoacott cagggatotg oggtogaggo tgtggoccoa gggoggaggo tgggtgoao aggtggooot gagototigic cacagactag agcaggaaag gggggaaagg oggogataga ggitagcagg aatgittaat tatcaggago iggaaggigg igggitgtoc iiiccacaco cotocototg aggigtgggo gtgggcoagg gotoacoaga ggcocoagag icititolgg gcccgaatag ccaccilggc agigtgggcg gcggctgccc tggtgacggt gcccacagct gittlegggg sgagggagge ctotgeactg traggaacag aggeagetet agtttggtte etgteatete tgggacaggg aaaceteeag agcatottoc toatoacago gotgagogtt gotogotact gggtggtggc catggotgog gggocaggoa occacototo ggagggga ggtgtgtgt gtgcgccttt gcctgctgcg tttccccagc aggtactggc tgggggccta ccagctgcag acaggoggg ogcatgoaca tataccotgg cattoaggot gtgootogoc otgoocoaco taccacoast ottgaocaac geaacggegg eggeaggaca geagggtegt ggecegetet gteegeatee tggtggette ettetteete tgetggttte occipicae lactigotig geacacagea alagoigeet caaccotigig cigiactgie tectgaggeg ggagecegg gggagiaaag taactotoco agtoacaegg clagtgagca geaggictgg gactoegeag ectoegetet tteeteet lggglggtgc tggctttcat ggtgcccttg ggcgtcatca ccaccagcta cctgctgctg ctggccttcc tgcagcggc ggacacccat getgattece tgeetetatg ceacetecea ggeecettge tittgggeece aagggaacae titttgeaga acciticagae acciticgici teaaccigge tetggeggae etgggacigg eacteactet eccettitigg geageegagi eggadtgga ctttcadtgg cccttcggag gtgccctctg caagatggtt ctgacggcca ctgtcctcaa cgtctatgcc scagtgigot gagtgotgat gatgotocga tgootgicaa attootagoo otgaggotoa tggttgooot ggootatggg citetegegg ccattegcit gcteggaaat itggeggtge tetgggtact gagtaactgt g ∞ ggagag ∞ ctgglphacaatetetga tgecetgega tgeceaeaet caataettet geeteteeae eeaeattett etgggeeaat geeteeggag cacatgratt ctctcattgc atcatgccac tcctgtgaag cagacttacc tgaaaatttt aagcaagaaa acaggcttag

iaagcaggta ggcaggcggt gggtcgcaag caaccccgg gagagccgcc cttctaccct gctcaccaac ctggacaga

Homo sapiens

sapiens

SALDFHWPFG GALCKMVLTA TVLNVYASIF LITALSVARY WVVAMAAGPG

AIGELGNLAV LWVLSNCARR APGPPSDTFV FNLALADLGL ALTLPFWAAE

Coupled Receptor 359

H7TBA62

G Protein-

190742

Homo

igicotocot goccoaaatg caaagcocag agtatcaatt tgagtgtoag agoaootgga ttoacagott tacotocago aaattaottt gittigittig ittgagacag agtotogito tgtogoccag gotggagtgo agtggtgtga totoagotoa otgoaacoto ogottocogg ccatticig cittogcaag aatacciagg aaaacticoc taagggitot aggciaaiga atcagaggic agtgcccate iciciolgia gagaataaac ctctggatta tocacaaatt gtcttgacct tttatoccag ttocacotoc agtteagtat ggaacaaaag gattegttgo sticaagoga ticicoigoc teagectece gagtagotgg gactacagge tecegetace atgeetggee aattittigt aattittaat agogattaa agaggggagg gggctgggag aacaggctgc aggtagagcc agaaaagcag agactccaga aagtggtgct occicitigi accicacigi tetcaacigi aaaaiggget actaaagati taacagigaa atatacigii agciattati ciigitigti gggggaggc gggggctcag atcagagctg gatgtgacaa agcttaagtc tttatttgga gatgggaaag aagaggatct gggaggaaa taagegtgca getgggagat ggggatgggg aaccatgtet cagetggaat ggttgtatat getetgaagt gacaccegg gignagggeg caagetgaac acactectet ttetgagate caceaagigt aggateetig agteetgggg itgocaggig igggggatig ciggaattic cagcaccigc caggcocigg gigiaaaacc ciggigciga cgggagigco stgotggagt tacaggogtg agocacogca cooggtogag ctattattot tacacootgt gtaaaatgga gacagagaga grepticic corctanate aggattigaa agaagtgaag ahaatgacaa gteaaagaca tgggtggggt gaagggaggt seggrataat gaaagtotoa cataaagaac toagaggttg goocotaago ocotottgaa ggtgtgttot ooaggacage gaagotgoc otototgoca ggotgcagtg cootcaggga aaaagtotga totttgatoc ocaactotgg gtgtggtgaa cottaccate teagtggtga ceactgaaae ttgetgeetg cagaggeete agetgeaaaa getgdagtte eettgaaggg gagacagag titcaccata tiggecagge tggicteaaa etectgacet etagtgatet geceaecteg geeteecaaa ocaccocc accteagage agggtatece tigtettet coggtateaa ggecaaaaat gecagettee eetgteetea agaaaggagg gaggcagagg gaagatgagg tagagctc MPTLNTSASP PTFFWANASG GSVLSADDAP MPVKFLALRL MVALAYGLVG gettectett tegeteetet atteggatge ateaatgata aaggitagee ateagaagga tittetagga ggeageeeet ENSP00000201

geteatett tateactgig etetteteea teateateig ggiggiggg ateteeatge teetgagagg eaaccogeag ticeagegae tecetegec atacitegea tegiggicae aatteigeta etettageat tietetteet eatgegaaag ateeaagaet geageeagig antgrocto cocaccoago tectettect ectgagtgre etggggetet teggactege tittgeette ateategage teaateaaca gtpactoto atcatgacca gaggtatgat gttfgtgaat atgacaccot gocagotoaa tgtggacttt gttgtactoc tggtotatgt gitoggggi igigicicci icicolggac gacaaticig igcatigota tiggitgcag totgitgcaa atcattatig ocactgagia nacigococo graegetaci trotettigg ggitetetti geretetgit teteatgeet ettageteat geeteeaate tagigaaget agottocaa giggagaaco aggagototo cagagoooga gacagigaig gagotgagga ggaigiagoa ttaacitoa ragetetigea ttetetacag ategigiaga caggagigee etttacaagg caatgeetige coegteacag cetaceaaca atgiacaagg actgcatcga gtccactgga gactattitc tictctgtga cgccgagggg ccatggggca tcattctgga itggiactec catteageeg cagacigitg ateceacaea agagigitte ateceacagg ctaaactaag eeceeageas etetteetg atggeectea cattettegt etecaaagee acettetgtg geeegtgtga gaactggaag eageatggaa gococagig ggacgaccog giogicigoa tigototiggi caccaacgoa igggittico igotigotigia catogitocot THESLEWARI ATLAVWAAAA LVTVPTAVFG VEGEVCGVRL CLLRFPSRYW VASFFLCWF PNHVVTLWGV LVKFDLVPWN STFYTIQTYV FPVTTCLAHS LGAYQLQRVV LAFMVPLGVI TTSYLLLLAF LQRRQRRRQD SRVVARSVRI NSCLNPVLYC LLRREPRQAL AGTFRDLRLR LWPQGGGWVQ QVALKQ

NM_018654

Coupled Receptor

GPRCSD

G Protein-

190743

Homo sapiens	Homo sapiens					Homo sapiens
۵.	∢	:			•	ρ,
Batgcaggag gagtataa MYKDCIESTG DYFLLCDAEG PWGIILESLA ILGIVVTILL LLAFLFLMRK IQDCSQWNVL PTQLLFLLSV LGLFGLAFAF IIELNQQTAP VRYFLFGVLF ÄLCFSCLLAH ASNLVKLVRG CVSFSWTTIL CIAIGCSLLQ IIIATEYVTL IMTRGMMFVN MTPCQLNVDF VVLLVYVLFU MALTFFVSKA TFCGPCENWK ÕHGRLIFITV LFSIIIWVVW ISMILRGNPQ FQRQPQWDDP VVCIALVTNA	WVFLLLYIVP ELCILYRSCR QECPLQGNAC PVTAYQHSFQ VENQELSRAR DSDGAEEDVA LTSYGTPIQP QTVDPTQECF IPQAKLSPQQ DAGGV CEEECagging eggaacctcc cigaagagtg ccciggicac agcacctig aagacagcca tiggicatigg ggacccaacc agaactigc ciggigagcca ggatgccat ccacaaagcc tiggigatigt gccigggaact gccicitic cigticccag	gggcctgggc ccagggccat groccaccg gctgcagcca aggctcaac ccctglact acaacctgtg tgaccgctct ggggcctgggg cccaggggcgt ggacggctct gggggggggg	ticacitytig cicigcigot gaccitigta gaggicatca toatacaga giggcigato alcaccitig ticggggcag tigggggag gegggaggggggggggggggggggggggggg	cittégret etteraegte ateccegagg teteccaggt gaccaagte ageccagage aaagetacca gggggacatg facceacce ggggggggggggggggggggggggggggggg	accecegaa gacegoaaga actotcaggi cittagaaac coctacegte eggactgagt cagcegrego gaggagaggo egggagaggo egggagaggo egggagaggo egggaggococ egggagaggococ cagactoc cotococte gcaggococ aacategococ cocactococ etotogoca gregitaggi eggitacat egtgococa cocactococ agtgittegi eggitaggi eggitacat estocagoca aatagtgit eggettegi eggtgega egcaaccoca ectocococ aggatcacot oggoegicac actocagoca aatagtgit tegggaga ticotgoaac otcaagagoc tocagococ	gaictigcte ctetiging accanging ectaniani acatticine tunitana annanana ana MGTQPEPGLG ARMAIHKALV MCLGLPLFLF PGAWAQGHVP PGCSQGLNPL YYNLCDRSGA WGIVLEAVAG AGIVTTFVLT ILL VASLPFV QDTKKRSLLG TQVFFLLGTL GLECLVFACV VKPDFSTCAS RRFLFGVLFA ICFSCLAAHV FALNFLARKN HGPRGWVIFT VALLLTLVEV INTEWLITTLVRGSGEGGP
NP_061124.1	NM_018653					NP_061123.2
190743 G Protein- Coupled Receptor GPRC5D	G Protein- Coupled Receptor	CARCOC.				G Protein- Coupled Receptor GPRC5C
190743	190744					190744
619	620					621

III-PRATANS QVMGSANSTL RAEDMYSAQS HQAATPPKDG KNSQVFRNPY VWD QGNSSAGWAV ASPCAVANND FVMALIYVML LLLGAFLGAW PALCGRYKRW RKHGVFVLLT TATSVAIWVV WIVMYTYGNK QHNSPTWDDP TLAIALAANA WAFVEFYVIP EVSQVTKSSP EQSYQGDMYP TRĞVGYETIL KEQKGQSMFV ENKAFSMDEP VAAKRPVSPY SGYNGQLLTS VYQPTEMALM HKVPSEGAYD

Ношо sapiens

sapiens Homo

NP 067647.1

Coupled Receptor

CGR7

G Protein-

190745

623

OLOKI YLONN KITSISIYAF RGLNSLTKLY LSHNRITFIK PGVFEDLHRL EWLIEDNHL PPLIFKDLKE LSQLNLSYNP IQKIQANQFD YLVKLKSLSL EGIEISNIOO RMFRPLMNLS ORKSMDSKGO KTYAPSFIWV EMWPLQEMPP ELMKPDLFTY PCEMSLISOS TRLNSYS /AIFLGINLA AFIIIVFSYG SMFYSVHQSA ITATEIRNQV KKEMILAKRF FFIVFTDALC VRPGKCRTIT VLILIWITGF IVAFIPLSNK EFFKNYYGTN GVCFPLHSED TESIGAQIYS WIPIEVVKFIL SLLQVEIPGT ITSWVVIFIL PINSALNPIL YTLTTRPFKE MIFRFWYNYR **JDCGNQADED NCGDNNGWSM QFDKYFASYY KMTSQYPFEA ETPECLVGSV** VQCLCQGLE LDCDETNLRA VPSVSSNVTA MSLQWNLIRK LPPDCFKNYH MISGSVFFYI-LIFGKYFSHG GGQDVKCSLG YFPCGNITKC LPQLLHCNGV HYFKKFQYC GYAPHVRSCK PNTDGISSLE NILASIIQRV FVWVVSAVTC SRISPPTFYG LNSLILLVLM NNVLTRLPDK PLCOHMPRLH WLDLEGNHIH SYNKHAQLWM ESTHCQLVGS LAILSTEVSV LLLTFLTLEK YICIVYPFRC NERNLTFISC SNLTVLVMRK NKINHLNENT FAPLOKLDEL DLGSNKIENI FGNIFVICMR PYIRSENKLY AMSIISLCCA DCLMGIYLFV IGGFDLKFRG

eccaatatoc tittgaggca gaaacacctg aatgittggt eggitetgtg ccagtgcaat gtettigcca aggictggag ettgactgtg caccogtit accigaiaaa ccicicigic aacacaigcc aagaciacai iggciggacc tigaaggcaa ccaiatccai aattiaagaa ccagaaact ggatgaattg gatttaggaa gtaataagat tgaaaatctt ccaccgctta tattcaagga cctgaaggag ctgtcacaa tagocttac taaactgrat otcagrcata acagaataac ottootgaag oogggrigttt ttgaagatot toacagacta gaatggorga gcatgogca gotgiggatg gagagtacto attgicagot tgiaggatot ttggocatto tgiccacaga agtatcagtt ttacigitaa itgaaaccaa titacgagot gitocatogg titoticaaa igtgactgoa atgicactic agiggaacti aataagaaag citoctocig ittgetteaa gaattateat gatetteaga agetgtaeet geaaaaeaat aagattaeat eeateteeat etatgettte agaggaetga gaatottic ctataatoca atocagaaaa ttoaagoaaa coaatttgat tatottgtoa aactoaagto totoagoota gaagggattg acatettog cagotetaaa ccaaacactg atggaattto atototagag aatotottgg caagcattat toagagagta tttgtotgg catagtitt ttectatgga agcatgtitt atagtgttea teaaagtgee ataacageaa etgaaataeg gaateaagtt aaaaaagag gratcige agtiaccige titiggaaaca tittigicat tigcatgega cettatatea ggtetgagaa caagetgtat gecatgicaa cattletet efgetgtgee gaetgettaa tgggaalata tttattegtg ateggagget ttgaeetaaa gtttegtgga gaataeaala catifotgac atiggaaaaa tacatotgca tigictatoc tittagatgi gigagacotg gaaaaigcag aacaattaca gitotgatto iggragaaat accaggtacc ataacctett gggtagtgat tittattetg eccattaaca gtgetttgaa eccaattete tataetetga aaatticaaa tatocaacaa aggatgttia gacotottat gaatototot cacatatatt ttaagaaatt ocagtactgt gggtatgcao aatigaaga taatcacctc agtogaattt ccccaccaac attitatgga ctaaattctc ttattctctt agtoctgatg aataacgtoc cattiggat tactggitti atagtggctt tcattccatt gagcaataag gaattittca aaaactacta tggcaccaat ggagtatgct ccotottoa ttoagaagat acagaaagta ttggagocca gatttattoa gtggcaattt ttottggtat taatttggco gcatttatoa atgacatotg gitotgiott ottotacato tiaattitig gaaaatatti tiotoatggg ggtggacagg atgicaagtg otocottggo ittigacitt taiticctgc agtaaittaa ctgitttagt gatgaggaaa aacaaaatta aicacitaaa tgaaaatact titgcaccic gateetige caaaegitit itettialag tattiaciga tgeatiaige tggataecea tittigiagt gaaatiteti teaetgette agaacatatg ctccatcatt catctgggtg gaaatgtggc cactgcagga gatgccacct gagttaatga agcoggacct atticocot gigggaacat cacaaagigo tigootoago tootgcaotg taaoggigig gaogaotgog ggaatoaggo ccacaagacc atttaaagaa atgattcatc ggttttggta taactacaga caaagaaaat ctatggacag caaaggtcag cgatgaggac aactgtggag acaacaatgg atggtccatg caatttgaca aatatttgc cagttactac aaaatgactt tteacatae ecotgigaaa igteacigat iteicaatea aegagaetea atteetatte atga

NM 021634 Coupled Receptor G Protein-190745

Egatgct gggacagggg traattgct gaagcaagtg ctctcatcc cctagctct gctgatctag grggacaggg traattgct gaagcaagtg ctctcatcc ctagctcta accateaa ctctgagctg ttggaca gaagcaagtg acttcagcc ttacctctt agccatcaaa ctctgagctg ttggaca gaacttcc ctaggcctct ctggccaca attctggc gaagaaaga gaaggaatga tacactc ctagggcaa gacacagga cgatcaggc actagtaga gagacactc ctggcaggc gttgccc acgcgagaa gacacagga gacacagga gagacactc ctggcaggc attggaga gacaggaga gagacactc ctggcaggc antcttg aagagaagg gagtcc gagagaagg gagtcc gagagaagg gagtcc gagagaagg gagtcc gagagaagg gagtcc gagagaagg gagtcc gagagaaga gacatggct ctgcatgtc ctgcatgttc catctttcga atctgctggc aactgcagga acatgagag gagtcc gatagaaga gacatggctc ttgaagtgc gacacagaagc gagacaa tgcaggccaa tgaaggaaga gacatgagga acatgagga acatgaggaca acatgagga atgaggaca acatgaggaca acatgagga acatgagga acatgaggaca acatgagaca ac	gictgggggt gggggatgct gggacagggg tcaattgcct gaagcaagtg ctctaiccc cctagctcct gctgatctag ttggggtcc agaggaatgc geggaaagg ttgggacc atttcgcc traccgtctt agccatcaaa ctctgagctg gaagaatga gaggaatga ggagatgca ggaggatgc actttcacct ctgggccaca attcctgcc gagagaaaga gaggaatga ggtgagcacc ttcttcactc ctagggccat gtggagagc tgcaggcac ctccttctg ccaataggca tagatgagg ggtgagcacc ttcttcactc ctaggaggg gtcaggaa ggcaaggt ccaatagga acagacacc tcttcactc ctaggaggc gtcaggaa ggcaaggt ccaatagga acagacacc tcttgaggc acctgcaca tgcagtgag gtgaagagg gtcaggaa ggcaaggt ccaatagga acagacacc ttgaaggc ttgaagtgc gtgaagagc gtgagagcc ggcaggagc gtgagagag gtcaggaa ggcaaggc tggaggcc ggagagagc gtgagagag gtgaaagaa gacaaagagg agcaggcc ttgaaggc ggaagagc agaaggagc ggaagagc ggaagaagc gaaagagc ggaagaagc ggaagaagc ggaaagaa gaaaagag gaaagaaga gaaaagag gaaagaag	egictgggggg gggggatgg transport gaagcaagtg transport gaagcaagtg ctctcatcc cctagctct gcgatctag tigggggt ggggatgga ggagaaggg transport gaagcaagtg cttctcgcc traccgtctt agccatcaa ctcgagctg gagatagtga ggagatgga ggagatgga ggggatagga ggggatggaca attctggcc attctggc gagagaaga ggagatggg gagataggacaca attctggc gagagaaga ggagagggggggaggaggaggaggaggag		AX147756
Egatgct gggacagggg tcaattgct gaagcaagtg ctctcaicot gtggggga gggacacggg tcaattgct gaagcaagtg ctctaicot gtggggag ggaacttc ctgggccac attcctgct ttggaca ggaacttic ctgggcctct ctgggccac attcctgc ctcactc ctaggccat gtggacacg ctccttcg ctgtgtgcc acgcgagca gccacagga cogtcagc actccttcg ctgtggat aaggaaggg gtccaggata gagcaaggt ccaatggtc attactgcc agatgca gaacttcc gaatggct attactgcc agatggat gaacttcgc gaaggatc gaaaggga gccaaggatc gaaagggc agaggaccaa tgcagggc agaggaccaa tgcaggga acaaaggg attactgcc tttgtaggc aaggga gaaagga ccacaatga gaacatgg gaaaggga atcaaggga atcaaggga atcaagtgc agagtga gaaaggga atcaaggga atcaagtgc acacaatgc caccaatgg tttataggc acaggaga atcaggaga atcaggaga atcaggaga atcaggaga atcaggaga atcaggaga atcaggaga atgagagg agacttcact accagggca atgatagg atcagt cacaagtgg atcagcaca acagccacaga tcaagtgg atgatagca acagccacaga acaagtgg atgatagca atgataggg atgatagc acacaatca a ggttcagc acacaatca a ggttcagca acagccaaga acaagatgg atgatagca acagccacaga acaagtgg atgatagca atgataggg agattcact taccagggca atgatagga atgatagg	gictgggggt ggggatgct gggacagggg tcaattgct gaagcaagtg ctctcatcd ttggggstcc agagtggga ggaaaggc actttgaaac ttctctgcc ttaccgtctt ggagtagtga cgatgtgaca gggaacttc ctgggcctct ctgggccaca attcctggc ggtgagcac ctcttcactc ctagggcat grgaaggc tgcagcaca attcctggc ggtgagcac ttctcactc ctagggcat gggaagtg cggaggat cagtgaa accgcaca tgccaggat agagaaggt ccgttcagca accaggat ttgaagtcg ctgggagtc gtgggatcg gtcaggata gagcaaggt ccaatgat ttgaagtcg ctgggagtc gtgggatcg ataacctca gcaatggttc ttgraggtc ctggagtc gggaagtcg ggaaataca gcaagaaga gacaagagt ttgraggca agaatggcg agaaataca gcaaagaag ttctaagga ggaaataca gcaaagaag ctctcaagga ggaaagga cacaatcaa ggaaaggac tcttcaagga cacaagagg caaggaagg caaggaagg caaggaagg		AX147756	GPCR Ls190748 AX147756 GPCR Ls190748 CAC39548.1
ggatgct gggacagggg toaatt gtggtga ggagaaaggc acttig tggaca ggagaatticc ctgggcc tcactc ctagggccat gtggtaga igtgaca acgcagaca gcaca cagtgat aaggatggg gtcag gagtcc gtggggatcg ataactt aatcttg agcattgcc agtagaa iaggagg ggaaataca gcaaa aagggc catcggtaagg cactaa taagcca tgaggaatag cactaa taagcca caccaatcaa ggtgtc tgcagc accaatcaa ggtgtc tgcagc accaatcaa ggtgtc tgcagc accaatcaa ggtgtc TAVLASLIIA TNTLVA LLTDQLSSPS RPTQKT FRYLKIMSGF VAGAC V FHPHFVLTLS CVGFF AG GYRSPRTPSD FKAI	gictgggggt gggggaaggg teaath tiggggctoc agagtgggga ggagaaaggc acttig gagalaaggc acttig gagalaaggc cattig gagalaaggc cattig gagacticc ctgggcc gggagactoc ticticacte ctagggcac ticticacte ctagggcac ticticacte ctaggacagc ggagagacag gggagagcag gggaaggc ggggaggg tittgaagtc gtggaagtc gtgggagtc gtggaactt tittgaagtcg ctgggagtc gtgggagtc gtgaaataca gaaat tittgaagtcg cagaatgagg gtgaaataca gaaat tocgagtggg aggaagcca tgaggaaga cactaa tottcaagta gcggaaggc tgcttgatgg cagaaggac gcttgatgg cagaaggac gattggtat cacaagaaggc acacaatcaa ggtgc tottgtggat caacagcagc acacaatcaa ggtgc tottgtggat caacagcagc acacaatcaa ggtgc tottgtggat caacagcagc acacaatca ggtgc tottgtggat caacagcagc acacaatca gagaaggg ttccatgtc cgaagtgg mESSFSFGVI LAVLASLIIA TNTLVA DTLIGVAISG LLTDQLSSPS RPTQKT FDRYLAIKQP FRYLKIMSGF VAGACCYKAEHAGAMAG GYRSPRTPSD FKAI		AX147756	GPCR Ls190748 AX147756 GPCR Ls190748 CAC39548.1
- 乳のほけない ほい 野路 思る 背 と ぬれ 口 ザ 4 人	giclggggt ge tiggggrtcc ag gagalagtga ce ggtgagcac th ggttgagcag gg actgcacaa tg tttgaagtcg cig tttgaagtcg cig tttcaagta gcg tcttcaagta gcg gcgagtggat caa actccaaatg ag MESSFSFGV DTLIGVAIS FDRYLAIKC YKGQCSFF		AX147756	GPCR Ls190748 AX147756 GPCR Ls190748 CAC39548.1

natiatitit taaaaaaaat tittaaaaag gittitigag acagaticit gcicigicac ccaggcigga gigcaglagc atgatcaggg

ateactgeaa ectetgeete etgggtteaa gegattettg tgeetaagee aeetgageag etgggattge aggtgeatge

angeattt geceantatt traeattigtt aetgeteaga ggratteett rattatgigg tragearagg trataetttig etgaegatte gotgggatt ataggoacaa gacaccacaa taattattgo ctgtatgtca attattattt taaaatattg ttgtatttac ttaatgtct

aagacagggt attgccgtgt tggccagact ggtctcaaac tcctgggctg aaacaatcct cccgccttgg cctcccaaag caccatgcct ggctaattit ggtattitta gtagagatga ggttttgcca ttttggtcag gctggaattt tttttttt taattitgal

sapiens

AACWLPYGCA CLAPAARAAE AEAAVTWVAY SAFAAHPFLY GLLORPVRLA

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GRLSRRALP GPVRACTPQA WHPRALLQCL QRPPEGPAVG PSEAPEQTPE

AGGRSPAYO GPPESSLS

egecgeatge etgragtece agetaetegg gaggetgagg caggggaatt gettgaacee gggaggegga gttttgecag eetggeaaca gageaagaet etgtetaaaa agaaaaaaaa attttttgt ttgagaeage alettgetet gteteceagg etggagegra aaggagate tettietgea tegacagaag tteetgeate ettteattea gagagacaga ggagaaagag tagteteatg tttteeteaa gaccaagat gaatagcaat acaattgctt ccaaaatggg tteettetee caateagatt etgragetet teaccaaagg gaacatgttg ictacaggia ctogocacca cacciggata attaaaaaat tattictgia gagatgaagt ctoactgigi tgcccagcci gggigicaal aagagatgg tgaagagact gcatgattaa actagataga cctggtatac agtcactgaa ctagtagatg tcaataatta ttattttaa gggocaatg attotagttt cagagtottg gaaggatgaa ggtagtgaat gtgaacotgg attittitog gaatggtaca toottgocat natigotigig tottatagaa otoaacatao tggggtottig aagattigtia ototgatigst ggoogttigg gtgotggoot tottagtigaa actectiag agccaggaga tiagccaagt cactggccat tototiaggg gittitgotg titgctgggc tocatatict ctgitcacas rgocatotot gaotitottig tgggtgtgat otocaticot tigtacatoc otoacaegot gitogaatgg gattitggaa aggaaatotg acateatte ttggaatteg tgateecagt catettagte gettatttea acatgaatat ttattggage etgtggaage gtgateatet ggaagactac acattttagg tatgtgatta gaaaacatac ttgtcagaat tgtctggctg gattaatttg ctaatttgac cttcttcatc grattings cteactactg actatetgit atgracagea tetgrafata acattgreet cateagetat gategatace tgreagtete tectettit gtatecatig tgteacaage gettteaaaa ggettiettg aaaatattit gtataaaaaa geaaceteta eeateaeaa icagteggte agtatettet taaagacaat ttteteaect etgtaaattt tagteteaat eteaectaaa tgaateaggt etgeoettta ctigocoti ticatictac caacagatot gcactitigaa gicaaiggia aattactoca gigaataata gcagtataat atgactigal gaaagtatg gettgteeca tttetteetg ttetetttt etagetteea eateagette etttttgag aacatataga agaagaagg gctataatg ctaggaatg ctttggreat titagettit gtggtggaca aaaacettag acategaagt agttattit ttettaaett ittigatgtg atgecagata ctaatageae aateaattta teactaagea etegtgttae titageattt titatgteet tagtagettt tetectite attitatice teageaacag grectaaate agittggtat agaatigeat ittggetiea giggiteaat teetitigtea gatcagtgg gtgggtgagg tagggtttga gttggcaaga gcagggaacg ggcatgtgcc caggtgagct cctgtgtgtg atattitig taaacitgta gicataatag tactatatic ticttagicc teacetette etigicitit agatettaat iteatgetga facaaaaat eegittigi titeitteta igiteeatge ataatacagi ettaagigaa titeiettii taatittat egiaatagaa iggicotcag igaagitatt itggaggooc iggiggicac aggatcagaa ggcaagggat aggcagtggt caccaatggi iggicaggag atogagacca tootggocaa catggigaaa ococatotgi actaaaatac aaacaagtag otggitgigg ccagattit ataticctaa tcccagtaag gaagaaagcg tagtgtggga gaggagagag ctgatgacig cagtictcaa aaatttitat tigitggoog ggcatggtgg otcacgootg aaatoccago acittgggag gocaaggtgg goggatcatg graatgeaat catageteae tgeageetgg aacteettgg eteaageaat eetgetgeet tggeeteeea agtatgtggg ocitatocag titgaaaate attocctaaa geatgeaata ggaaaaagaa ceteetgget gggaetgeee aaetetgtie aglaggige caaagecate etggaetgae tgetgtetet tecaacatet giggaeacte atteagaggt agactatett

Histamine H4 Receptor 190774

NM_021624

	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
acatitiati agtitiggita tgittigice tittaaaaca titticittig agatgggggg citigototigi tgoccaegea ggagtgeagt gggatgeagt gggatgetet cagotoactg cagocotgac tgoctagget ocagoatet tottaogica gcotocagag tagotgggac egocactaaa aatititiaa attgitigot itottaaagt giticitigoc igicitigic acaaaattic attiticica tagitaatit catotococg giaagatiti attgitigiti cititataac tugocagite tiacaccgit tigigatiti catgiticit agaactita aacottiaac ticaaacat aaaatacaag tottitaagt acatgagge tiagaaatgi acataatgit tatatacact attgoctiac attaaagtoc aatatgagaa atacatgiti aacattcaat aataattita aaaatttgag aaataaacto toataaatgo aaaaaaaaa	MPDTNSTINL SLSTRVTLAF FMSL VAFAIM LGNALVILAF VVDKNLRHRS SYFFLNLAIS DFFVGVISIP LYIPHTLFEW DFGKEICVFW LTTDYLLCTA SVYNIVLISY DRYLSVSNAV SYRTQHTGVL KIVTLMYAVW VLAFL VNGPM ILVSESWKDE GSECEPGFFS EWYLLAITSF LEFVIPVILV AYFNMNIYWS LWKRDHLSRC QSHPGLTAVS SNICGHSFRG RLSSRRSLSA STEVPASFHS ERQRRKSSLM FSSRTKMNSN TIASKMGSFS QSDSVALHQR EHVELLRARR LAKSLAILLG VFAVCWAPYS LFTIVLSFYS SATGPKSVWY RIAFWLQWFN SFVNPLLYPL CHKRFQKAFL KIFCIKKOPL PSOHSRSVSS	cagiccag gagcagacaa galggagaca aattoctoto tatoto ttoctggala toatcacita totggattt geagtcacot tggatto eggatgacac acacagtcac caccatcagt catt ottoatggta gagaaggcaa tgggaggaca ttggcottto tgtt oggaagtgto ttoctgattg coctcattgo totggaccgc cgtgag cotggcaag aaggtgatca ttgggcoctg acag tacctggtaa aacggggaca gtagoctgca cttttaactt tgttgcca tgttgacggt gagaggcatc atcoggttca tgttgcca tgttgacggt gagaggcatc atcoggttca tgttgccat atcaggtggt ggcocttata gccacagtca tgct tgtgcccat atcaggtggt ggcocttata gccacagtca ccggga gatgtgaca atcaggtggt cttcttcaac cgggga gatgtgaca accitcagc cttcttcaac ccggga gatgtgatc cacgcccttc cogccagtct cagctacca attctacttt accttctgca gaggtggagt cccagctc cagcttcgtc teaccttgag ttaggctgag aaaa aaaaaagcct ttgtgtccc tgatttgggg agaataaaca	<u>a</u>	atggaaacca actictocat toctorgaat gaaactgagg aggtgctoco tgagootgot ggocacacog tictgtggat ctictcatig A H ctagtocacg gagtcacctt tgfcttoggg gtcotgggca atgggctigt gatctggglg gctggattcc ggatgacacg
8 · 题 · 音 · 题 · 语	NP_067637.2 M S3 C3 C4 C9 C9 C4 C4 C4 C5 C5 C5 C6 C6 C6 C7 C7 C7 C7 C7 C7 C7 C7 C7 C7 C7 C7 C7	NM_002029 CC TO		NM_002030 at
	Histamine H4 Receptor	Formyl Peptide Receptor 1 (FPR1)	Formyl Peptide Receptor 1 (FPR.1)	Formyl Peptide Receptor-like 2
		190823	,	190824
		630	631	632

sapiens Ното

gggtgatgac gggactotgg attitcacca tagtecttac ettaccaaat tteatettet ggactacaat aagtactacg aatggggaca ceacticat tatiggette aeggigeeta igtocateat cacagietge tatgggatea tegetgeeaa aatteacaga aaceacatga acagotgoot caacccaatt ototacgtot ttatgggtog taacttocaa gaaagactga ttogotottt gcocactagt ttggagagg acagicaac accaicigit accigaacci ggecciagci gacticicti icagigecai eciaceatic egaaiggici eagiegeca iggragictg geteraaagag atgitgitaa atggeraaata caaaateatt ettgteetga ttaacceaae aageteetig geettittta catacigiai titcaactit gcaticiggg gigacacigc igiagagagg tigaacgigi tcattaccat ggccaaggic titcigatcc gagagaaaaa tggccttttg cgtcattoct atgtaagtta gttcatgtta tgatagacat caacctgttt gtcagtgtot acctgatcac taaatocag cogtooctta ogtgtottog otgotgtggt ggottottto ttoatotgit ggttooctta tgaactaatt ggoattotaa cootgactga ggicootgac teageceaga ecageaacae acacaecaet tetgetteae eteetgagga gaeggagtta catcattgot of ggacogot graftigigi cotgoatoca gootgggood agaaccatog caccatgagi of ggoodaga caagcaatgt ga NP_002021.2 Formyl Peptide Receptor-like 2 (FPRL2) 190824

633

(FPRL2)

IVPMSIITVC YGIIAAKIHR NHMIKSSRPL RVFAAVVASF FICWFPYELJ GILMAVWLKE MILINGKYKII LVLINPTSSL AFFNSCLNPI LYVFMGRNFQ ERLIRSLPTS LERALTEVPD FIIVLTLPN FIFWTTISTT NGDTYCIFNF AFWGDTAVER LNVFITMAKV FLILHFIIGF JHVMIDINLF VSVYLITIIA LDRCICVLHP AWAQNHRTMS LAKRVMTGLW AGFRMIRI'VN TICYLNLALA DFSFSAILPF RMVSVAMREK WPFASFLCKL METNFSIPLN ETEEVLPEPA GHTVLWIFSL LVHGVTFVFG VLGNGLVIWV SAQTSNIHIT SASPPEETEL OAM

NM 013447

EMR2 Hormone

190948

634

Receptor

gicateacet acatgggget gagegtetet etgetgtgee tecteetgge ggeceteact titeteetgt gtaaageeat eeagaaeae ctgggagcat ggccagaatg gatgtggtca ctgggccacc acaggctgca gcacaatagg caccagagac accagcacca secgeceggg etggeaaceg atteeggggt ecceaatgg eccaaacaat aeegtetgtg aagatgtgga egagtgeagt saatgaatge aceteeggae aaaacceatg ecacagetee acceaetgee teaacaaegt gggeagetat cagtgeeget vataacacca tecagageat ettacaggeg cfggatgage tgetggagge ecetggggae etggagaece tgeceegett ctggagteca cagecagaeg etttecegat tettegacaa agtecaggae etgggeagag aetacaagee aggettggee eggagaceggg acagecetgt eccaeteact etttecectg etgeteetge eggeagetea getggaaeca tgggaggeeg ctgctggaac acagaggga gctacgactg cgtgtgcagc ccaggatatg agcctgtttc tggggcaaaa acattcaags itgagagega gaacacgtgt caagatgtgg acgaatgtca gcagaaccca aggctctgta aaagctacgg cacctgegt nacaccotog gcagotacae gtgccagtge etgectgget teaageteaa acctgaggae cegaagetet geacagatgt cogggeage atcagtgiga cagetecace gtetgettea acacegtggg tteatacage tgeegetgee geceaggetg saagcccaga cacggaatec cgaataacca aaaggacaet gtetgtgaag atatgaettt etecaeetgg accegeeee scagcagcac tgtgtggcca gtcacctgct ggatggccta gaggatgtcc tcagaggcct gagcaagaac ctttccaatg gacagaatc aggcagtgat gcagcicgac tggaatcagg cacagaaatc tggtgaccca ggcccttctg tggtggcc atcaccaccc ccatggagac ttgtgacgac atcaacgagt gtgcaacact gtcgaaagtg tcatgcggaa aattctcgga getettgaa etteagttat eetgeaggea cagaattgte eetggaggtg cagaageaag tagacaggag tgteaeettg gictocatt ccagggatgg gcaagtigct ggctgaggco cototggtoo tggaacotga gaagcagatg ottotgcatg ctgoogttg cacccacctg agcagctttg cogtoctcat ggcocactac gatgtgoagg aggaggatcc ogtgotgact igacacacca gggcttgctg caggacggct ccccatcct gctctcagat gtgatctctg cctttctgag caacaacgac ggtggtgccc tcaggactcc tcgtgtgtca atgccaccgc ctgtcgctgc aatccagggt tcagctcttt ttctgagatc egictlicic gictlicieg caticigigi ciggcigaci cigeceggag cigaaaccca ggactccagg ggcigtgcc occaaaacc teagetecce agtiacette acettetece acegiteagt gatecegaga cagaaggige teigigieti

socigiacet ettecteaet geaeggaace tgaeggtggi caactaetea ageateaaca gatteatgaa gaageteatg

tecciging getaeggagi eccagetgig acagiggesa ittergeage etceaggest cacettaig gaacacette

icacaaggig cigigotoca icatogoogg tacotigoac tatototaco iggocacoti cacotggatg otgotggagg

geaceteae tgeatetgea getetegete tgeetettee tggeceaect ectetteete gtggeaattg ateaaacegg

Homo

Homo sapiens

NP_038475.1

EMR2 Hormone

190948

635

Receptor

cegetigetigg etecaaccag aaaaggatt tatatgggge ttecttiggae etgetigege eafettiett grgaatttag ttecttiet ggggaetiet tiggagget tectecete aataggaag tgtocaccet eeggaacaca aggatgetigg cattataage gacagetigg gaggatetig gaggatetig gaggatetig gaggatetig gaggatetigg aggatetigg aggggaet aggatgetigg gaggatetigg aggatetigg aggatetigg aggatetigg agggatetigg aggatetigg aggatetic eatiggaget tragggga

gecattetet cacatecegt geggteagga agecetteet gaactetgae tteagtiett getgeggttt etgeceattt tttleatate MGGRVFLVFL AFCVWLTLPG AETQDSRGCA RWCPQDSSCV NATACRCNPG GLLNFSYPAG TELSLEVQKQ VDRSVTLRQN QAVMQLDWNQ AQKSGDPGPS EPVSGAKTFK NESENTCQDV DECQQNPRLC KSYGTCVNTL GSYTCQCLPG VVGLVSIPGM GKLLAEAPLÝ LEPEKQMLÍH ÉTHQGLLQDG SPIĽLSDVIS AFLSNNDTQN LSSPVTFTFS HRSVIPRQKV LCVFWEHGQN GCGHWATTGC NNOKDTVCE DMTFSTWTPP PGVHSQTLSR FFDKVQDLGR DYKPGLANNT TALKPEDPKL CTDVNECTSG ONPCHSSTHC LINIVGSYQCR CRPGWQPIPG SPNGPNNTVC EDVDECSSGQ HQCDSSTVCF NTVGSYSCRC RPGWKPRHGI QVGPAARVMA YLFTIINSLQ GVFIFLVYCL LSQQVREQYG KWSKGIRKLK QSILQALDE LLEAPGDLET LPRLQQHCVA SHLLDGLEDV LRGLSKNLSN IIAGTLHYLY LATFTWMLLE ALYLFLTARN LTVVNYSSIN RFMKKLMFV GYGVPAVTVA ISAASRPHLY GTPSRCWLOP EKGFIWGFLG PVCAIFSVNL STIGTRD TST ICRCTHLSSF AVLMAHYD VQ EEDPVL TVIT YMGLSVSLLC ELLAALTFLL CKAIONTSTS LHLOLSLCEF LAHLLFLVAI DOTGHKVLCS ALFLVTLWIL KNRLSSINSE VSTLRNTRML AFKATAQLFI LGCTWCLGII SSFSEIITT PMETCDDINE CATLSKVSCG KFSDCWNTEG SYDCVCSPGY TESEMHTLSS SAKADTSKPS TVN

gocattotot cacatocogt goggicagga agocottot gaactotgao ticagitott gotgoggit otgoccatti tittoata ototgacago tgogaggica tototgotot ggottitoto caagcagaac aagtgggggo totggaaagg taagggaco tcagtggoca ccattataot tigcatotti cotgagaagt gagagttgaa agggaagcag gaaggoccai ggtcagattg aaggaaggao tittagitt cittittitti tittgaaat ggagtotogo totgicatto aggotggag gcagtggtgo gatotcago cactgcagoo tocacttoot gggttcacat gattotog cotcagoot caaglagot gaagacaca gcacatgoca

NM_000752

Leukotriene B4 Receptor BLT1

Ношо

Homo sapiens

Leukotriene B4 Receptor BLT1

190955

637

191039 Trace Amine

Homo sapiens

NM 022049

Coupled Receptor 88 (GPR88)

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grecticataa itotgaccae actogitgge aatotgatag itatigitie tatateacae itoaaacaae itoataccoe aacaaatigg	cicaticati ccatggccac tgrggactit citctggggt grctggtcat gccttacagt atggtgagat ctgctgagca ctgttggtat	inggagad icincigna dancacaca agcaccgaca trangctgag cicagccicc attiticcati igiciticat ciccatigac	egetactate eteretere iccacteaga tataaageca agatgaatat ettegitatt tereteatea tetteattag tiggagigte	octgot gttt ttgcatttgg aatgatottt ciggagotaa acttcaaagg cgotgaagag atatattaca aacatgttca ctgcagagga	ggitgototg tottottag caaaatatot ggggtactga cotttatgac ttottttat atacotggat ctattatgtt atgrgtotat	tacagaatat atcitatogo taaagaacag goaagattaa tragigatgo caatoagaag otocaaatig gatiggaaat	gaaaaaitgga atticacaaa gcaaagaaag gaaagcigtg aagacaitgg ggaitgigai gggagtitic ctaataigct	getgecettt ettatetgi acagteatgg accetttet teactacatt attecaceta etttgaatga tergitgatt tegitigget	actignacte tacattaat ceaniggitt atgeatitt etateetigg titagaaaag eactgaagat gatgetigtti ggraaaatti	tocaaaaaga ttcatccagg tgtaaattat ttttggaatt gagttcatag	MMPFCHNIIN ISCVKNNWSN DVRASLYSLM VLIILTTLVG NLIVIVSISH	FKQLHTPTNW LIHSMATVDF LLGCLVMPYS MVRSAEHCWY FGEVFCKIHT	STDIMLSSAS IFHLSFISID RYYAVCDPLR YKAKMNILVI CVMIFISWSV PAVFAFGMIF	LELNFKGAEE IYYKHVHCRG GCSVFFSKIS GVLTFMTSFY IPGSIMLCVY	YRIYLIAKEQ ARLISDANQK LQIGLEMKNG ISQSKERKAV KTLGIVMGVF	LICWCPFFIC TVMDPFLHYI IPPTLNDVLI WFGYLNSTFN PMVYAFFYPW	The state of the s
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Receptor I (IAI)									-		191039 Trace Amine	Receptor 1 (TA1)					
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ocogaagic attitggacg gocacotgat tittacocit igtitotgig tittagagga atociaaagi caaaacaca gagactigaa

saactigoaa actiggogtit taaaataaco ggitaatita titicoacaca gittigititt gaaaaagago titoalaatig tataacoot

ccactitica tegicitata tatgaagege citigagigig catgaaccaa aggaaataac attgaagaag gaaaacaata

sapiens

NP_071332.1

Coupled Receptor

G Protein-

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88 (GPR88

Ношо coaggacait aggaccacit gitgiacate tgaataatta tggaagtigg gacatgitaa ggaaaacaaa tatgitcate accaacaate tagacaagg atattitact tettecagae accagaagaa atggeettea attatttgaa aagagacaea gagacaeete tggetaeeta accatgact gcatagctaa tattagctgc tattgcatgc toctagatgc tagaacttat tgggcatgtg gtatactgaa gcgatacccg gagitettee tetettgace aattiatgag aaageteeca gitgggacti tateteacaa gtggaateae agteaagaeg gateaataat itegitegot cagcaaagoo agotetecto titlagget taaacaagoo acacettaga aagcaacact giittatet agticatata iattaccacg acatttaaca tcaatattgt atatgttgaa ggaggtataa taaactcagt catatatagt gaacagttca aalgggaaag aggigigoco accagtaiga gitgocatta agacotcaag cocittaito itaaaagggi titlaataaa gictitotca aalgaggiag natettagee agtgagaaaa aaaattatti tatgeteett tittitegea etettaagae tgaaaattgg egttgagtgt tatagtgaaa uttitecagt itgataatig afggteagag ecageactgg aattitgaaa acaaataagg igaitateta ititaggiae egtiteaeal gragaaagt attttagaaa gtaacctgtc tttgatgatg cttctcttac catttagttt ttgtatatta coctggggca gtgaagcoct grantight gotaagaaga ataagteett etgittiete titaaeattt aaaatatete aatgeaeatg atataattaa aeaetaataa atcaccttat caaattaaaa tgggaagaaa gtaattttaa taattttaa taatcatatg tcagcattct gactacttac cacatcaaai iticialiage atgeacacit gitgelacce teatitigia accaatitat tigecitiatg aatgigatig cagettigaa cattetgiae gtictaaaa catattatti gaggttigic ataticatci tiggttiact aaattiacti agaaatatti gaaatgcaaa atigigigaa agotgicati trattaatot atocottitig igoatgoaco attictotot tactaacagi ticatotigti cacattitico tigaticaaa ctgggcccaa acagoctcag ttaactgcat aattcaggaa caaaaccago ttgctttgtt gcacgcctgg gcaatttcag MTNSSSTSTS STTGGSLLLL CEBEESWAGR RIPVSLLYSG LAIGGTLANG tattaaagtt cagaaaaaa aaaaaaaaaa aaaaaaaa aaaaaaa

iatogatogo taccagaaga ccaccaggoo atttaaaaca tocaaccoca aaaatotott gggggctaag attotototg ttgtcatotg geaticaig ticttactot citigoctaa catgatictg accaacaggo agoogagaga caagaatgig aagaaatgot citicottaa itgictitiga cigcactgct gaaaatacte igtictatgt gaaagagage actotgtggt taacticctt aaatgcatge ctggatecgi aggaccactg agaactttig tgtgtcaagt tacctccgtc atattttatt tcacaatgta tatcagtatt tcattcctgg gactgataac catctattt ttteettige aagteettea gaaatteett gataagtatg etgaagtgee eeaattetge aacateietg teeeaggaea actgototac actgrootgt tittigtigg actiateaca aatggootgg ogatgaggat titotiicaa atcoggagta aatcaaact attatttt ettaagaaca cagteattte tgatettete atgattetga etttteeatt eaaaattett agtgatgeea aaetgggaae atcagagitic ggictagict ggcatgaaat agtaaattac atcigicaag tcaittictg gattaattic ttaattgita tigtatgita caaagtttt cattaicatt getgiattet ttatttgttt tgtteettte cattttgeee gaatteetta caeeetgage caaaeeeggg acactcatt acaaaagaac tgtaccggtc atacgtaaga acgagggtg taggtaaagt ccccaggaaa aaggtgaacg gcogtogaca acoteacete tgegeetggg aacaecagte tgtgeaecag agactacaaa ateaeecagg tectetteee ggotgoaata actactactt actggataca ttcaaaccct ccagaatcaa cagttatcag gtaaccaaca agaaatgcaa ALYORRHTAG MLALSWALAL GLVLLLPPWA PRPGAAPPRI HYPALLAAAA PPADWDGAGG SYRLLRGGLL GLGLTVSLLS HCLVALNRYL LITRAPATYQ MVIYLVSSFR KLQTTSNAFI VNGCAADLSV CALWMPQEAV LGLLPTGSAE LLAQTALLLH CYLGIVRRVR VSVKRVSVLN FHLLHQLPGC AAAAAAFPGA /WVSLASGFS LPVPWGVHAA SWLLCCALSA LNPLLYTWRN EEFRRSVRS\ OHAPGPGGAA HPAQAQPLPP ALHPRRAQRR LSGLSVLLLC CVFLLATOPL LPGVGDAAAA AVAATAVPAV SQAQLGTRAA GQHW

sapiens

Homo

191168 P2Y12 Platelet NM_022788 ADP Receptor

ataggaaaaa agaacaggat ggtggtgacc caaatgaaga gactccaatg taaacaaatt aactaaggaa atatttcaat

logatergea gitgragect gigetgiggt giggateatt teactggiag etgteattee gatgaeette tigateaeat eaaceaaeag gaecaaeaga teageetgte tegaeeteae eagtteggat gaacteaata etattaagtg giacaaeetg attitgaetg eaachaettt etgeeteeee tiggigatag tgaeaettig etataeeaeg attateeae etetgaeeea tggaetgeaa aetgaeaget geettaagea

gaagcacga aggciaacca ttotgctact cettgcattt tacgtatgtt ttttaccett ccatatettg agggtcatte ggatcgaate togcetgett teaatcagtt gttocattga gaatcagate catgaagctt acategttte tagaccatta getgetetga acacetttgg

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			ctottigigi tcagaactog itaaagcaaa gogotaagta aaaatattaa otgacgaaga agcaactaag itaataataa tgactctaaa gaaacagaag attacaaaag caattitcat ttacotticc agtatgaaaa gctatottaa aatatagaaa actaatctaa actgragotg	
191168	P2Y12 Platelet ADP Receptor	NP_073625.1	MOAVDNITSA PGNTSLCTRD YKITQVLFPL LYTVLFFVGL ITNGLAMRIF FQIRSKSNFI IFLKNITVISD LLMILTFPFK ILSDAKLGTG PLRTFVCQVT SVIFYFTMYI SISFLGLITI DRYQKTTRPF KTSNPKNLLG AKILSVVIWA FMFLLSLPNM ILTNRQPRDK NVKKCSFLKS EFGLVWHEIV NYCQVIFWI NFLIVIVCYT LITKELYRSY VPTPGVGYVD BYKYNNYGYJFI INAJEFICEV BEHTEA DIDXT I SOTTO DAYBOA	Homo sapiens
			TAENTLEYVK ESTLWLTSLN ACLDPETYFF LCKSFRNSLI SMLKCPNSAT SLSQDNRKKE QBGGDPNEET PM	
191193	Trace Amine Receptor 3 (TA3)	AF380189	¥	Homo sapiens
			concacute anacacuge acacacua anacanteg angegrege iggeotyge tgacnteng grgggagtea etgigangee concacute anacaacua geografica ethicageas geografica geografica in ggggacagti actgiaaati ecatacatgi titgacacat cettergiti tgeitentia geografice tgitgataga tacatigetg tlactgatee tetgacetat ecaaceaagi tractgitgte aqtitteaggg atatgeatgt tettiteetg gtiettitet geografica gettiteeat ethicaca geografica agtitteaggg atatgeatig tiettiteetg gtiettitet geografica gettiteeat ethicaca gaacaaca aagaagaat	
•			tgaggaatta gragtigoto taacotgigt aggaggotgo caggotocac tgaatcaaaa ctgggtocta ctttgitto tictatiott tatacocaat gtogocatgg tgittatata cagtaagata titttggtgg ccaagcatca ggotaggaag atagaaagta	· ·
			cognostical agencial to concegada guacangga angagungan anangagngan ganaggongo cananoning ggantigota tegoragonit totigicatot tegoraccal accitogitga igoagigati gatgottata tgantittat anotocotoci taigititatig agattitaga tegorgigat cagotataga occottgati tatgottici titaccanig gittigggang	
191193	. • -	AAK71240.1	taa P	lomo
	receptor 3 (1A3)		CIVLSWFFS	sapiens
			VAMVETYSKI FLVAKHQARK IESTASQAQS SSESYKERVA KRERKAAKTL GIAMAAFLVS WLPYLVDAVI DAYMNFITPP YVYEILVWCV YYNSAMNPLI YAFFYQWFGK AIKLIVSGKV LRTDSSTTNL FSEEVETD	
961161	G Protein- Coupled Receptor	AF411109	catocca A attttc	Homo sapiens
•	GPR80	· · .	aaaalgagac ctiggaagag cagcaccatc attatgciga acctggccig cacagatcig ctgtatciga ccagccicoc cticcigait cactaciatg ccagtggcga aaactggatc titggagat tcatgtgtaa gittatccgc itcagcitcc atticaacci	

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Homo	Homo		Homo sapiens	Homo sapiens
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taacctgita ctatatgigg tegtcagcga caactticag caggctgtct gctcaacagt gagatgcaaa gtaagcggga accttgagca agcaaagaaa attagtact caaacaaccc ttga MNEPLDYLAN ASDFPDYAAA FGNCTDENIP-LKMHYLPVIY GIIFLVGFPG NAVVISTYIF KMRPWKSSTI IMLNLACTDL LYLTSLPFLI HYYASGENWI FGDFMCKFIR FSFHFNLYSS ILFLTCFSIF RYCVIIHPMS CFSIHKTRCA VVACAVVWII SEVAVIPMITF-LITSTNRTNR SACLDLTSSD ELNTIKWYNL ILTATIFCLP LVIVITCYTT IIHTLTHGLQ TDSCLKQKAR RLJILLLAF YVCFLPFHIL RVIRIESRLL SISCSIENQI HFAYTVSGPI AALNITFGNI ILYXVXVSDNFO OAVCSTVRCK VSGNI FOAKK ISVSNNP	AT	agraccetra geacegage descriptor greatgige ceatetigia togetgoege egococagae acetgreage gigtegitgit greetgetet gaggocotgic extactgetg agralettiga aaggaaagit etgiggette thaittaatga alggigacte tiggitgitgit cagacattig attenteae tgeagogigg etgatittiit taiteatiggi teetigiggg teeagtetigg ecciptiggi cagacattig attenteae aggatetige acagacage egantetigg ecciptiggi ecciptiggi excepting exceptiggi exceptig	MDPTTPAWGT ESTTVNGNDQ ALLLCGKET LIPVFLILFI ALVGLVGNGF VLWLLGFRMR RNAFSVYVLS LAGADFLFLC FQINCLVYL SNFFCSISIN FPSFFTTVMT CAYLAGLSML STVSTERCLS VLWPIWYRCR RPRHLSAVVC VLLWALSLLL SILEGKFCGF LFSDGDSGWC QTFDFTTAAW LIFLFMVLCG SSLALLVRIL CGSRGLPLTR LYLTILLTVL VFLLCGLPFG IQWFLLWIW KDSDVLFCHI HPVSVVLSSL NSSANPITYF FVGSFRKQWR LQQPILKLAL QRALQDIAEV DHSEGCFROG TPEMSRSSLV	tcatatacht gacattottt ttogaggoaa agtittagat acacitgtgg catittocot gcatatgtgt gcaaatgott gtgcotgaag atcittgctt ttotgcoagg ttgcagactt gccactagag ctgggattgg tcattgtgac attgcogotc atggagtoca gtgaagcagg actoagggca atgotgotca cactatggga agaataactg tagaicatot tgagaaaggo agacittgig taatotott gottacaaaa
CAC51133.1	AY042214		AAK91805.1	LG94359
G Protein- Coupled Receptor GPR80	MrgX2 G Protein-Coupled Receptor		MrgX2 G Protein-Coupled Receptor	G Protein- Coupled Receptor Ls191222
191196	191218		191218	191222
647	648		649	650

toatalacht gacatictit tiegaggeaa agtittagat acacitigigg catiticct geatatgigt geaaatgett gigectgaag A alcitigett tietgecagg tigeagacti gecaclagag etgggattigg teatigigae attgecgete atggagteea gigaageag acteagggea atcagggea atgetgetea caclatggga agaataactig tagaicatet typigaaaggea atgetgetea caclatggga agaataactig tagaicatet typigaaaggaea atgetgetea caclatggga tigaataactig atataatat gacaataate tecacagcig glacatatt gecaaatigig grageataga taggateaa getatgaagt aaatgageat gecaaatga atgaattigg ettecatiga attecatat tigectitiga augeaaatat gaageaaata aaggecaagga tegecaatga gecaaggag caagtatgaga teoctectea caclecagga tgatgactet gggeaaggag acatteacet etacagtagg tgetgecaaag attagecaga gtgtgeaaat gacaacctgg atgecegte aagtgaagaa aattagagaga atattgggat caaagctgaa ggetageaaa attiteagag acttegteaa aatteggaga caaagctgaa tgttgectgg tittacatgt gaagtettgt ggttetecaa tgaaaaaggt eggegea tegageaaagaaaagga taaaaggaga aaatteegaa aattagggat caaaacttg tittacatgt gaagtettgt ggttetecaa tgaaaaaggt egtgectae tecaaacatt gtetgectgg tittacatgt gaagtettgt ggttetecaa tgaaaaaggt egtgectae

sapiens

Homo

sapiens

Homo

cattocagtt gagatatico acticcitit caaagcacat agigciccia acagggoco agigagitti gitgitgoat aaaaggcagi icitigiaaat attatgecaa caaccagaac aaatatgatt cocagtaggg agagaatcag gagtaggatg gecaaggagt aaattgagga aatgacagag aaggatcaca tagcagactc ttaatccccc ggatgatttc acaacaggtg tgttcaggtt

OTLAMIHSIE MINNSTILPG VKLGYEIYDT CTEVTVAMAA TLRFLSKFNC ENSP00000199 Coupled Receptor

G Protein-

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Ls191222

EANNVCIAFK EVLPAFLSDN TIEVRINRTL KKILLEAQVN VIVVFLRQFH VFDLFNKAIE AEILSDKIRF PSFLRTVPSD FHQIKAMAHL IQKSGWNWIG IITTDDDYGR LALNTFIIQA SRETVEFKCD YSSYMPRVKA VIGSGYSEIT MAVSRMLNLQ LMPQVGYEST

CHILPSDSHK LLHEYAMHLS ACAYVKDTDL RLIHSIQLAV FALGYAIRDL VININKMWIAS DNWSTATKIT TIPNVKKIGK VVGFAFRRGN ISSFHSFLON

COARDCONPN AFOPWELLGV LKNVTFTDGW NSFHFDAHGD LNTGYDVVLW KEINGHMTVT KMAEYDLQND VFIIPDQETK NEFRNLKQIQ SKCSKECSPG

OMKKTTRSQH ICCYECQNCP ENHYTNQTDM PHCLLCNNKT HWAPVRSTMC

EKEVEYLNW NDSLAILLI LSLLGIIFVL VVGIFTRNL NTPVVKSSGG LRVCYVILLC HELNFASTSF FIGEPODFTC KTROTMFGVS FTLCISCILT KSLKILLAFS FDPKLOKFLK CLYRPILIF TCTGIQVVIC TLWLIFAAPT VEVNVSLPRV IILECEEGSI LAFGTMLGYI

ALLAFICFIF AFKGKYENYN EAKFITFGML IYFIAWITFI PIYATTFGKY VPAVEUVII

SNYGILYCT FIPKCYVIIC KQEINTKSAF LKMIYSYSSH SVSSI

iticitgage taggaaaggt ggitggetta eggeacagta gagagettee agggetgget ggegtgggat accegtacea NM 032571

-agaaatgea gggaceattg ettetteeag geetetgett tetgetgage etetttggag etgtgaetea gaaaaceaaa aetteetgtg ctaagtgccc cccaaatgct tectgtgtca ataacactca ctgcacctgc aaccatggat atacttctgg atctgggcag aaactattca cattoccott ggagacatgt aacgacatta atgaatgtac accaccotat agtgtatatt gtggatttaa cgotgtgtgt tacaatgtog

octeticagg acaccactic cicaaagaca accgagggca ggaaagagti gcaaaagati giggacaaai tigagicaci icticaccaat cagactitat ggagaacaga agggagacaa gaaatcicat ccacagctac cactaticte egggatgtgg aatogaaagt totagaaact goottgaaag atocagaaca aaaagtootg aaaatooaaa aogatagtgt agotattgaa aaggaagttt ctactgtcaa tgtgtcccag gatatagact gcattctggg aatgaacaat tcagtaattc caatgagaac

Receptor EMR3 Mucin-Like Containing

EGF-Like

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Module-

tgeagtgae ateateeagg gagacacaca aggteceagt gecattgeet trateteata trettetett ggaaacatea taaatgeaae acteaagega ttacagacaa ttgetetgaa gaaagaaaga catteaactt gaaegtecaa atgaacteaa tggacateeg

ittittgaa gagatggata agaaagatca agtgtatctg aactctcagg ttgtgagtgc tgctattgga cocaaaagga acgtgtctct icacagggca gggcagccag tggtccaggg atggctgctt cctgatacac gtgaacaaga gtcacaccat gtgtaattgc stocaagiot gigacgotga cittocagoa ogigaagaig accoccagta coaaaaaggi citotgigio taciggaaga egicaccigi ccagciticge igiccigaig geocigacea gecaggagga ggatecegig cigacigica teaectaegi

xotcactgca totgcagoto togototgco tottootggo coacotooto ttootogtgg ggattgatog aactgaacoc aaggtgotgt sofocatoat ogcoggigot tigoaciate totacotggo ogcottoaco iggaigotgo iggagggigi goaoctotto otoaotgoac gaacctgac agtggtcaac tactcaagca tcaatagact catgaagtgg atcatgttcc cagtcggcta tggcgttccc gggotgago gictototgo igigoctoot cotggoggoo otcactitto tootgigaa agcoatocag aacacoagoa

segaticate tegagiticc tiggoccagi cigigocati tictorgoga attragrati gittatotig gictitigga tittgaaaag agaactitoc teceteaata gigaagigte aaccatecag aacacaagga tgetggetit caaageaaca geteagetet catectggg etgeacatgg tgretggget tgetacaggt gggtecaget geecaggtea tggeetaect etteaecate solgigacia igaccattic igcagocico igaccicaco ittaigaan igcigatega igcigacico acciggacca

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atcaacagoc tocaaggott ottoatotto tiggiotact gootootoag coagoaggto cagaaacaat atcaaaagtg gittagaaga atogtaaaat caaaatotga gottagaaga atogtaaaat caaaatotga gottagagaca tacacaottt coagoaagat gggtootgac tocaaaacoca gtgaagggga tgutticoa ggacaagtga agagaaaata taaaactag aatattoaac tocatatgga aaatcatato catggatoto tttggoatta tgaagaatga agotaaggaa aagggaatto attaaacata toatoottga agaggaagta atcacottt acttoccaag cettiette tocacaatag gotocoaa aateteret aaattecatt tottoaaa aaaaaa	MÖGPLILIPGL CFLISLFGAV TÖKTKTSCÄK CPPNASCVNN THCTCNHGYT SGSGQKLFTF PLETCNDINE CTPPYSVYCG FNAVCYNVEG SFYCQCVPGY	RLHSGNEQFS NSNENTCQDT TSSKTTEGRK ELQKIVDKFE SLLTNQTLWR TEGRQEISST ATTILRDVES KVLETALKDP EQKVLKIQND SVAIETQAIT DNCSEERKTF NLNVQMNSMD IRCSDIIQGD TQGPSAIAFI SYSSLGNIIN ATFFEEMDKK	DQVYLNSQVV SAAIGPKRNV SLSKSVTLTF QHVKMTPSTK KVFCVYWKST GQGSQWSRDG CFLIHVNKSH TMCNCSHLSS FAVLMALTSQ EEDPVLTVIT YVGLSVSLLCLLLAALTFLL CKAIQNTSTS LHLOLSLCLF LAHLLFLVGI	DRTEPKVLCS IIAGALHYLY LAAFTWMLLE GVHLELTARN LTVVNYYSSIN RLMKWIMFPV GYGVPAVTVA ISAASWPHLY GTADRCWLHL DQGFMWSFLG PVCAIFSANL VLFILVFWIL KRKLSSLNSE VSTIONTRML AFKATAOLFI	LGCTWCLGLL QVGPAAQVMA YLFTIINSLQ GFFIFLVYCL LSQQVQKQYQ KWFREIVKSK SESETYTLSS KMGPDSKPSE GDVFPGQVKR KY	KHAYICLAAI WAYASFWTTM PLVGLGDYVP EPFGTSCTLD WWLAQASVGG QVFLNILFF CLLLPTAVIV FSYVKIIAKV KSSSKEVAHF DSRIHSSHVL EMKLTKVAML ICAGFI IAWI PVAVVSVWA FGPPDSPIO I SVVDTI IAK SAANNDIN	QVIDYKFACC QTGGLKATKK KSLEGFRLHT VTTVRKSSAV LEIHEEV	abyganyan veeebyeey eegaagege toeggoogog gagagege igoocagag goegoegeg igogaaagoo gegoogogogogogogogogogogogogogogogog	caeecegaee eeceeee eegeeeegc ceeegeceg graeegecc geeageeegc ceaageeeg eegcaegcca aaegcccega cceeegeeeg eegceegiega egcceigaag egaegceega batpategcg agegeege
	NP_115960.1					CAC21687.1	NM 001407	101-100 TAIN	
	193511 EGF-Like Module-	Containing Mucin-Like Receptor EMR3				G Protein-Coupled Receptor	Cadherin EGE	LAG Seven-Pass	G-1ype Receptor 3 (CELSR3)
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GAITLOAPLD YEDQVTYTLA ITARDNGIPQ KADTTYVEVM VNDVNDNAPC

GLVTLALPLD YKQERYFKLV LTASDRALHD HCYVHINITD ANTHRPVFQS

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FVASHYTGLV SEDAPPFTSV LQISATDRDA HANGRVQYTF QNGEDGDGDI

TIEPTSGIVR TVRRLDREAV SVYELTAYAV DRGVPPLRTP VSIOVMVODV

NDNAPVFPAE EFEVRVKENS IVGSVVAQIT AVDPDEGPNA HIMYQIVEGN

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> NP 001398.1 LAG Seven-Pass G-Type Receptor Cadherin EGF (CELSR3)

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MKVGVLWLIS FFTFTDGHGG FLGKNDDIKT KKELIVNKKK HLGPVEBYQL LLQVTYRDSK EKRDLRNFLK LLKPPLLWSH GLRIIRAKA TTDCNSLNGV LQCTCEDSYT WFPSCLDPQ NCYTHTAGAL PSCECHLNNL SQSVNFCERT KIWGTFKINE RFTNDLLNSS SAIYSKYANG IEIQLKKAYE RIQGFESVQV TQFRMSLLSP KLECNGTI	argagricot goaacticac acargocaco tutgigotta tiggiatoco aggarlagag aaagoccati totgggitgg citococoto citicoatgi atgiagtigg aargicaalog tgglottoat ogtaaggaeg gaacgoagoc tgoacgotoc gatgiaocto tutocatgi atgiaggaego tagingaalog tgglottoat ogtaaggaeg gaacgoagoc tgoacgotoc gatgiaocto tutocagaa tgottagaago cattgaoctg gocttaroca catocaccar goctaagato citigoctti totggittga ttoccagaag atiagottig aggoctgtot taccagaag tiotifaito atgocototo agocatigaa tocaccatoc tgotggocat ggocttigao cattagogo catotgogo aargotgoag tootaagaa tocacaataa goccaarate	gcalogiggo tgrggroego ggarocolor tituticos actgociotos cigalcaago ggotggoott orgocactoc aatgrooto organicola trigigocaco caggargiaa tgaagtiggo ciatgoaga actitigoca atgrigata tggictarot gccaticigo tggicatggo cgiggacgia atgricator cottgicola titorigaia atacgaacgg ttorgoaci gccttocaag tcagagggg ccaagggocit tggiaaccigi ggicacaca ttggitggi actogocit tatgigoca tratiggoci ctagtigta caccgcttig gaaacagoci tcatoccatt ggotgittig foatggiga catcaccit cigcigoci cigcigoci cigatgaga caccacati gaacagoci catcacata tocatacati tatgigagaca	MISCNETHAT FVLIGHGLE KAHFWVGFPL LSMYVVAMEG NCIVVFIVRT ERSLHAPMYL FLCMLAAIDL ALSTSTMPKI LALFWFDSRE ISFEACLTOM FFIHALSAIE STILLAMAFD RYVAICHPLR HAAVLNNTVT AQIGIVAVVR GSLFFFPLPL LIKRLAFCHS NVLSHSYCVH ODVMKI AYAD TI PNVVYCH TAIT VAGYON	MFISLSYFLI IRTVLQLPSK SERAKAFGTC VSHIGVVLAF YVPLIGLSVV HRFGNSLHPI VRVVMGDIYL LLPPVINPII YGAKTKQIRT RVLAMFKISC DKDLQAVGGK actitutica igutciccit gagtgaagga igaggaaati gaaagcagag tatgcaccit tiattaggag attcaaactg catcctactg gattagcctc aaaagtccta aaaagccta aaatacaaag acatccatct gacagatcac taaggagag actitetitit ctettinga anaetticce
NP_079324.1	NM_030774		NP_110401.1	NM_032787
194319 G Protein- Coupled Receptor FLJ22684	Olfactory Receptor, Family 51, Subfamily E, Member 2		Olfactory Receptor, Family 51, Subfamily E, Member 2	194743 FLJ14454
194319	194431		194431	194743
099			662	663

atfaacatoc egaateecat gtgeaetgeg attgeegeet tactgeacta ttttetgtta gtgaeattta eetggaaege aeteagegel

gcacagetet attacettet aataaggace atgaagcete tteeteggea ttteattett tteateteat taattggatg gggagteeea

aatcactiga catattatoc aacgtiggal gigcactgic igitactggi ciggototica cagtiatati toagatigto accaggaaag

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ccaataagaa citgcagaca agtgatggtg acatcaataa tattgacitti gacaataatg acatacccag gacagacacc

Homo

sapiens

Homo

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aaagotttaa toootggaaa ggocaogaac aatgaatooa titoalgoat ottgitggaa caoototgoo gaactitiaa acaaatootg ggeaacate eteatigiai teactataat aagatecagg aaaaaacag teectgacat etatateige aaectggetg tggetgattt gaataaagag tttgettate aaactgecag tgrggtggat acagteatee teeetteeat gattgggatt atetgtteaa cagggetgg scateateae atecetggat aettgraace aatttgeetg tagtgeeate atgaetgtaa tgagtgtgga eaggraettt geeetegtee eggeeggeegg cagggttege gaggeaceea egetectaaa aagageacga egeaceegat geteggattg gatgaagtge getocacata gitggaatge citticitat teaceaatgg geogggggg gagagtgggt gittggggggg extetetgea

KKVSSMKKI VSTLSVAVVF GITWILAYLM LVNDDSRIV FSYJFCLFNT TQGLQJFILY

VRTKVFQSE ASKVLMLLSS IGRRKSLPSV TRPRLRVKMY NFLRSLPTLH

ERFRLLETSP STEEITLSES DNAKESI

DYRQEKICW LAIPEPNGVI KSPLLWSFIV PVTIILISNV VMFITISIKV LWKNNONLTS

YYLLIRIMK PLPRHFILFI SLIGWGVPAI VVAITVGVIY SQNGNNPQWE

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DAQTELQVL LNMTKNYTKT CGFVVYQNDK LFQSKTFTAK SDFSQKIISS

OKGTDGFLRC RCNHTTNFAV LMTFKKDYQY PKSLDILSNV GCALSVTGLA

TVIFQIVTR KVRKTSVTWV LVNLCISMLI FNLLFVFGIE NSNKNLQTSD SDINNIDFDN NDIPRTDTIN IPNPMCTAIA ALLHYFLLVT FTWNALSAAO

cogagta ctgaggaaat cacactotot gaaagtgaca atgcaaagga aagcatotag acagtaaaac ttacotgttg tggtotttt gagadante tgetggetgg caattecaga acceantggt gttataaaaa gteegetgtt gtggteatte ategtaeetg taaecattat ceteateage aatgitgita tgittattae aatetegate aaagtgetgi ggaagaataa eeagaaeetg aeaageacaa aaaaagttte haattettt acaagtaet ahaaaggaca caaagagaaa actttaeett eeagaacaaa atgaeteetg atgaacagtg tgtggggat agaattica cacaacatac aagagtacca tigitoctia tatogitaaa tottigigac acacttigac aaaaatgtag aacctataac gcatcagg ategretica getacatatt ergeetttte aacactacae agggattgea aattittate etgtacaetg ttagaacaaa tecatgaag aagattgita geacattate tgttgeagtt gittitggaa ttacetggat tetageatae etgatgetag ttaatgatga tattagataa aacotgitgi ttattattat teggeataat ggaettggta gittitetat titteaatag attigtaett gaataaggtg ateaceteg titgagitti atetgittet etecittati toccagicet eteagaaagi etiecteaat giattiiget eaggattaag scogagget gegtgtaaag atgtataatt tecteaggte attgecaaee ttacatgaae getttagget aetggaaaee ctatagtag tggctataac agtgggagtt atttattotc agaatggaaa taatocacag tgggaattag actacoggca sicticcag agigaagcit ccaaagigit gaigitgcia togiciaitg ggagaaggaa gicatigcci tcagigacgc Ecitigidg tathaactt tigacctctg
MASCRAWNLR VLVAVVCGLL TGILLGLGIW RIVIRIQRGK STSSSSTPTE

> FLJ14454 194743

NP_116176.1

CRNGGTWEN GRCICTEEWK GLRCTIANFC ENSTYMGFTF ARIPVGRYGP

SEQTOGKD TP NAGNPMAVRL CSLSLYGEIE LOKVTIGNON ENLETLEKOV

KKVAIVTVSQ LLDASEDAFQ RVAATANDDA LTTLIEGMET YSLSLGNOSV

JEPNIAIQSA NFSSENAVGP SNVRFSVQKG ASSSLVSSST FIHTINVDGLN

EDVTAPLNNÍ SSEVQILTSD ANKLTAENIT SATRVVGQIF NTSRNASPEA

NM_032503 Coupled Receptor SLT/MCH2 G Protein-194745

	Homo	Homo sapiens		•	Homo sapiens
aaccatticg actgacacgt tegagaacaa gglacaagac catocegatc aatttgggoc tttggggagc ttoctttatc ciggcattgc ctgictggg cractegaag gicarcaaat thaagacgg tettgagagt tegtgetting atttgacatc coctgacgat gracicting attgacatc coctgacgat gracicting attgacatc coctgacgat gracicting attgacata acaactttit tittocctor acccitigatt titggtggc atatttaat tittatgctat actifting attgacaaca gaataaggat gcagargct gcaatoccag tgaaccaaa cagaragtga tgaagtgac aaaaggagt gcagargct gcaatoccag tgaaccaaa cagaragtga tgaagtgac aaaaggagtgac tictatgrg tgaagtct tatoctgag gctactatg gcagacgat acatggga aactacaag tgaacaacga tgaaaattc cagaaacgtc tgoctcaaat ccaaagaaga gcgactgaga aggaaatcaa caatatggga aacatctga aatacacact ttagaaagga acatggatca agacatgat glotatctta ctggtattat tagaaagggc aggtaacgaatatgatat gcaaatgat glotatctta ctggtattat tagaaagggc aggtaacgaatatgatat gcaaatgat gaaaaggg taaacatga aatacaatga gcttaatatag ctaactgaa aaaaaaaaaa	MAPPHASCWN TSAELLNKSW NKEFAYQTAS VVDTVILPSM IGIICSTGLV GNILIVFTII P RSRKKTVPDI YICNLAVADL VHIVGMPFLI HQWARGGEWV FGGPLCTIIT SLDTCNQFAC SAIMTVMSVD RYFALVQPFR LTRWRTRYKT IRINLGLWAA SFILALPVWV YSKVIKFKDG VESCAFDLTS PDDVLWYTLY LTITTFFFPL PLILVCYILI LCYTWEMYQQ NKDARCCNPS VPKQXVMKLT KMVLVLVVVF ILSAAPYHVI QLVNLQMEQP TLAFYVGYYL SICLSYASSS INPFLYILLS GNFQKRLPQI QRRATEKEIN NMGNTLKSHF	ccacacaca aggacocgca toctgggtga tgaagtcaga cacgcagcag ctgggtgagt gctaacgctc agataagcat ctgggcatt gtggggactc cctgggctgc tctgcacocg gacacttgct ctgrococgc catgacaac gggtcggct googcatt gtggggacacc atctcocagg tgatgcogcc gctgctcatt gtgggcctttg tgctgggcgc actaggcaat ggggtcgccc tgtgtggttt ctgcttccac atgaagacct ggaagcccag cactgtttac cttttcaatt tggccgtggc tgatttoctc cttatgatct gcctgccttt tcggacagac tattactca gacgtagaca ctgggctttt gggggcattt cctgcccagt ggggctttc	acettegeca tgaacaeta triccacce getegeget egenteget tecegacaeg tatticaaag tegtccacce caccaegeg tgaacaeta triccacce getegeget egenteget geaccity gecociggi atcatege atcategaa accatetot gegtegeget gegeteget geaccity gegcociggi atcategaa accatetot gegtegaaa geeggeceg tototigiaaga getteataa gegeggaa aaccatetot gegtecaaaa geoeggeceg tototigiaaga getteataal geggtegec aatgategaa accatetot gegtegaaaa geoeggeceg catcateta tittgeteci tcaagatigi ttggagcctg aggeggegggaagge aggaggaagget eggtegaaaaggat eggtegaaaaaggat eggtegaaca geggegacca gaataaaaaaaaaaaaaaaaaaaaaaaaaaaaaa	attigggoott geacataaco deagottea concattaa cagcatgotg gatocoetag tgeutgegan conceptual attigggoott geacataaco deagottea concatgaa cagcatgotg gatocoetag tgtattattt tteagocco teottteeca aattictaaa cagcataa atetgeagot tgaaacoeta geagocagga cacteaaaa cacaaaagoc ggaagaaga cocaatticga acoteggiog caggaagtige ateagtiggg caaatagtig caaatiggt gaagtigge antigggatec caacaatagtig acottigga acticaaagocag totgatgggg acttagaatt aactegiget aaaggaggtegg aggettigaa aatteggaago acottictaa ttgeaagaaga tagagtggga acttagaactgc atectictea ttetgiegga aatgaaatte acacaactat acottitiggg gaggtteeag tt	MYNGSCCRIE GDTISQVMPP LLIVAFVLGA LGNGVALCGF CFHMKTWKPS TVYLFNLAVA DFLLMICLPF RTDYYLRRRH WAFGDIPCRV GLFTLAMNRA GSIVIEF TVÄVA ADDVJEVAVATED UTAAVARTISTID VAA GDIVCTT 31 VII CTTAVA
	NP_115892.1	NM_032554			NP_115943.1
	G Protein- Coupled Receptor SLT/MCH2	Chemokine Receptor FKSG80/GPR81			Chemokine Receptor FK SG80/GPR81
	194745	194756			194756

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MYNGSCCKIE GD I ISQVMPP LLIVAFVLGA LGNGVALCGF CFHMK I WKPS TVYLFNLAVA DFLLMICLPF RTDYYLRRRH WAFGDIPCRV GLFTLAMNRA GSIVFLTVVA ADRYFKVVHP HHAVNTISTR VAAGIVCTLW ALVILGTVYL LLENHLCVQE TAVSCESFIM ESANGWHDIM FQLEFFMPLG I ILFCSFKIV WSLRRRQQLA RQARMKKATR FIMVVAIVFI TCYLPSVSAR LYFLWTVPSS ACDPSVHGAL HITLSFTYMN SMLDPLVYYYF SSPSFPKFYN KLKICSI.KPK

Receptor FKSG80/GPR81

sapiens Ношо

OPGHSKTORP EEMPISNLGR RSCISVANSF OSOSDGOWDP HIVEWH

stcatggagt gtctgcacgg gacgtcctgg agagtcggac acgtaagcag cacagtgagg ccaccaacag cagcaaccg acattegeat ceteateget gtgaccagag teateteaca gateagegee gacaactaca agatecatgg agacceagt stettegtgt aetgegeett eetggaette ageteeggag aaggggtetg gtegaaceae ggetgtgege teaegagagg aaccicacc tactccgtct gccgctgcac teaectcace aactttgcca tectcatgca ggtggtcccg ctggaggtca

caacggitgi geigtggiti tecagtacai gitigecacg eteaacteee igeagggaet giteatatte etetticati gieteetgaa

cottoaagt tgaeggecaa ggeagtegee gtgetgetge ceateetggg tacetegtgg gtetttggeg tgettgetgt

géocrticea eteggacete algaatggga eceggecagg catggectee accaagetea gecettggga caagageage

tragaggig agagoogoot toaagoacaa aaccaaggio iggiogotoa ogagoagoto ogocogoaco tocaaogoga

cactotoccc acceptega cotetcagec gigigageog ggaggetecc aaccagecca ggetecete agaacacac

seccitgag ectecettea teacteagea teagaceeag egaggeeagg acaetegggg eeggteeege ageaeeagga

agacagotg tecteccotg tgaetetgge tgteggagea cactgeteag eccageagee tgatgeecag gecagegtgg

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ocotoctgo ettgeateea ecegtggget gagtgaette etegggggat teeeaggaea cagtggeetg aetgtgatgg

gggatgite agectetgig cettggtggg gettgggggae teagggecaa agaggtggit eaggteecea egeaecetea

caggogca ggcagotggg ggtgtgtggg gaagagcatg oggagtcccc agtgtctgaa tccactgagt ggtgagttcc

cacagoogg operagoogt ggtgtgtgt totgtaggtg gtgcoggogt gggccaacot gtgctgtgtc atcagttggg

Homo

sotgatagca ctogtggtgt ctgtgaaatg tgggtaagac attcaaacot ggttttgata otggaaacte tteetttaaa aetgtgaeca gatticati cagococtoc acacocotat giotigootig ittoagagig agitticiat ggagcotgig gocotitigo agocoacotg aaagaatgaa atgittagit tatagtagaa gaaagatgat gacactaagt tgtgaaaata tgitgigatt titatgaaat aaactcatgi aregeceate egeigigagi itigeoteti iggaececaa iteggeetia agaigecete etecetegig igecageete etiggilgi traatigit geogratica tetatatage taatattica agataagtaa tgaacaaaae etgtetaaae etttigitte caatgaatga aagtcatgca ctitatitat aggototatg tittggotto tgcagtactt ttattatota tacataattt ggocaaaaat aagaaattgg caatgracac ttggatattt ctccttattt agtttctagt gaaacaaatc aagtaaggaa ctatctttag tttagatgga attatttgtt atggigacag tgccacgggc cotgegtatg gecootgcaa cogtgototg gegggcacae etggetgotg caggocaagg getecgtet ggteaecatg agaccgaect gegetgagte eccaetgaec tggagaggga gggetggtga cagecgtgte tteggocac aggagotego cetetococg cagtgooteg tetocagete gaaagtegag ggcattitoc agggoacteo igaaaccagg tcacatggac cacagtgcca gatecteate aegeeggtga geaectagaa gtgagaacae tgtatteeta secotgoc caageogage tegagoogig ggogggagte gitgaetete caggtgaggg ogaeceetet geootgtoo geagggage ageatgtotg caggggtgaa cetttgetet totgteagge gaggeceagg etgeaecage caeetgeeac itococaga ggottocica iggotoacag goactotacg aagittotaa igggoagacc aegeggoagg tageacagtg gogggggtc cocicigcic acgigaagag cogciciggg cottgaggci gootgatggt gootgfgctt gggggagett totetette agggaaattt atggactcag actcagcccc agaggagatg ggataattgt tatggaccca tgtgtgggca stggettett aafgtaacte tteecetggt egeetggagt ggaceactea tetgeaggee teteetgeat ggggagggta gatocigig gaacacaggi tigggatoat agatgigaat taagacacca oogagataog ggotgigagg ticataotgi cognightea gigaagagic ceatgittag tatggactaa agreecatgi tiagecactg ecceaggete eegtgacee

HGVSARDVLE SRTRKQHSEA TNSSNRVFVY CAFLDFSSGE GVWSNHGCAL HIGDPSAFKL TAKAVAVLLP ILGTSWVFGV LAVNGCAVVF QYMFATLNSL IRGNLTYSVC RCTHLTNFAI LMQVVPLEVN IGILIAVTRV ISQISADNYK

> G Protein-194757

670

CAB82385.1 Coupled Receptor Ls194757

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Coupled Receptor

Ls194757

G Protein-

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QGLFIFILFHC LLNSEVRAAF KHKTKVWSLT SSSARTSNAK PFHSDLMNGT RPGMASTKLS PWDKSSHSAH RVDLSAV	ttágitcaag iccaggicga cactgottig gotgottggg tggiaggcaa igotgggoc gggactgicc ogggaggotc ttoccacag coctgoagg cacottiggg oggotgocci coagggggot gigiagogot gategocag coccaiggot acggggacactg cogotgcact ggcacticot agggaagaga gggacaacag tgioccaggo cocagiggog ggogotgotc	alaggecagg actgagagga geagtgtgge cacgtaggge coccageaca geoegaagag cagcatgget ceagcetgg cotttgectg extecaggta agggeoeggg ecagggegga gggeteateg eggeacactg ecgetecag eggeagatg teetgeaget ggeggtggge agtggecage acgeggacag agaggaagge ageageacec aeggegggaa geaggageece	angacticg aggracaggi agggggctgg gaagatagcc tgggagctgc agttggcacc aggggtccag tggttccacc ccagagcggg cagactggca aagagcaggg gaccagcca ggtgaggatgg agggggttgg aggggcttca ggatgctccc aggggggttgg aggggcttca ggatggcat gaatggtccagc ccgtgcacca gcaagaggtt ggcaagcagg gaagggaagg	egicaagus avgaggagus aggaicagu accoggoga ciciggiico acagoocigg caalgigggo aaigocagac Cogtgagcag cocagocago agiaggotca ggaagaagca gocagcaggt gggotgogca ggoggoggto ccaggogaig Cocagoocia poppocadott for or or or an accomplance a manacococa an accococa translance	gottgeberger Sergeriger er e	MIPNSI GEVP SPIPKGALGI. SLALASLIIT ANLLLALGIA GTAACAATCW LLLPEPTAGW AAHGSGIATL PGLWNQSRRG YWSCLLVYLA PNFSFLSLLA NLLLVHGERY MAVLRPLOPP GSIRLALLLT WAGPILFASL PALGWNHWTP	GANCSSQAIF PAPYLYLEVY GLLLPAVGAA AFLSVRVLAT AHRQLQDICR LERAVCRDEP SALARALTWR QARAQAGAMI. LFGLCWGPYV ATLILLSVLAY FORPPI GPGT 11 ST 1ST GSA 8A AVDXAMG I GDODVTA DW DODDV CDG	CGEEPPGTVP APALPTTQAA KAVSTWT CGEEPPGTVP APALPTTQAA KAVSTWT Caggeccag gatagagtaa teategggte cacageactg getagatgag tegegetett treatectaa tettatooc	afgitagcac agaacitigig tegcagtaga gagaggicag gcitcagagi cagcaagaac tegaiticaa actegaitig ággacccca ccittigata ggigacitat tcictgigag tcictgatci gcccictita aatgaggaag taaatcccac atggcaggg
	LG94710					553		AY042215	
	194858 G Protein- Coupled Receptor LS194858				G Protein-	Coupled Receptor LS194858		MrgX3 G	Protein-Coupled Receptor
,	194858				194858		. •	194878	
	671	٠.	•		672			673	

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aggacocca cetttigata gatgacitat tototigian gootlettia aatgaagsaag taaatocca atggcaggit
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ggittigigit gaaacgicaga atticatiaa aatogocgigo eggittiti tatgitigigi tototigigg tocagoctig totocico
totigigoci gocottigo atticatigo atticatica catogotigo ecatococa cattactic tietiggatiga aagtotiat tigicatigi catocico
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gaagotigi totocagagg gotogocaga gagagatiga ggigatiga ggitegaggat titotagga goticaaaat agcagaacc
tgaagotigi totocagaag gotogocaga acacococa cattactic tietigggot cottuaggca titaagagcaa igcigoccig
gaagotigi totocagaagi totocagaaagita agaacatogo citaacaggact tigagaagaa igcigoccig
coaccitiga caattatalig cattatigo cicagaaaatig

Ношо	sapiens	•				Homo	sapiens		
д						≺			
MDSTIPVLGT ELTPINGREE TPCYKQTLSF TGLTCIVSLV ALTGNAVVLW	LLGCRMRRNA VSIYILNLVA ADFLFLSGHI ICSPLRLINI RHPISKILSP VMTFPYFIGL	SMLSAISTER CLSILWPIWY HCRRPRYLSS VMCVLLWALS LLRSILEWMF	CDFLFSGADS VWCETSDFIT IAWLVFLCVV LCGSSLVLLV RLCGSRKMP	LTRLYVTILL TVLVFLLCGL PFGIQWALFS RIHLDWKVLF CHVHLVSIFL SALNSSANPI	IYFFVGSFRQ RQNRQNLKLV LQRALQDTPE VDEGGGWLPQ ETLELSGSRL EQ	tcaggtggag ccgcagcgcc tcgtgragtc ctgaatggag gcctggaagt gctctgtgct gttgaggtct gggcggcaga	ggatcaegta geacitagge agaaaatace cacegaagee getgeteagg etgeteagee cagecateat gttggeegea	ggcaggiact igccgicgta gacgctggcc gtggtgaaga aggcgatcca ggacacgaag ttgaagagca ggctgaaggt	gacacattig ecclettet aeticicles caagicotta occapitae tecapecaa pecacipate papagene
AAK91806.1				•		LG100657			
8 MrgX3 G	Protein-Coupled	Receptor				G Protein-	Coupled Receptor	GPCRB3	
194878						194903			

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attocctag caggcagtgg ggtccacacc accagccaag ttagacagat aagcagctgg gccgctgage tgatcatcac

aaacaggcca gcaccgtggt tttggaccca ggcgtggtag aatgtaggta cettggtgga aaacttgaag atgatgatta

goottgtgg gitococaaa gaagocatag aggotgocac tacotgotgc cagggagocc agcalaagaa agcacaggog cocootgot gacotcacca caggggtgtc taggtgocag gcaaacaggo cagcagtocc aagcagcago agcagcago

jogigitago igocagoago acceaagag igigoteacg caaagocaaa aacaceacag igogogggaa geaggicigg

tatteattt agagaaagag gitgaattea ggataegaet gettitgiag gagtigtgat gaeagetete taacagagga caeaecteag titigggg ggacgaatte tegetitigtg gtecaggetg gaatgeatet tggeteaetg caaceteege etectgggtt caagtgatte cotgenica gentenegag tagnigggat tanagnang ngenacan occapnaat tititatatt titiggtagag atggggtf ggcacacgc cacaacgccc ggctaactti tittgtatti tiagtagaga tggggttica ccatgtiggt caggctggtc tcgaactcc itgetetet gitettgace itgeatitet ggalggggaa igetgitti tietetgetg cagacaeget aglatetgia iteaggeeas geaccigia agaagecaga ggggceacae giaggggeec aagteaaagg acageteaca igiggaacag aaaacagaa cigiticaag gagciagcig tottiggcat gggcaacaga agggacagta ggacaagagg gcacaaaggg aacaatagci ictggcagcc cagtgactgg gccttggttc tggggcaggg cacatggggc ccaagggagg coctecetec accgtgcag ngagccagat gagcagagta ggaataggaa ataggggcct gcaagatact gggagaattg taccagggca gctagacta gglaggicitg cocacataco agagaggita ogatotgatg ggagcagoot gotocoaagg gagggcatig taacocotot otottgocag catticcatg aaccactitic otgagotgot gototgiggi tiototgagi ootgacooto tgaggacaga aggg acacteaaa geageagtga tggaaaceeg taaceacteg etggtgeeet teaagaeagt egetggaaca cacagaetta ctaaggett teagtigget aattettet ttettettt tttttgaga eagagttit etettgiege eeaggetiga gigeaatgg icotcaggi galocaccea cologgeote ceaaagigel gggaliacag gigigagoca cogegeoegg cotectitet socioggagi gotgggrago togotigoto cattgoocac toaccactot tottgaggaa ggtoocagoo coacaggga ictaggeata gigggatggg ggtagccggg agtggggcct gaggccacgc atticcicaa aatgccigtg ttaattacag cotgrocot acagagatgg tgaaaggaaa gaatgtggco cotggacaco aactaaggao otgagroott agotacotaa iccatgitgg ccaggctggt ctcgaactcc cgacctcaag tgatccaccc gcctcagcct cccaaagtgc tgggattaca citocotoag graccoacte itetitocea caaggeigge atetgiagag gietgaaagg gaaggeeaag aaggitoetg geafgagee accgeaceea giggetgaft etettgatea gaattetgte iggiageagg igteeteeaa cetgaageta scietiggaga cacacaggie ggitetgiat ggeteatgat cecatgaggg tittgeaaac eetagggagg acettaaeet caatottgg ctcactgcaa cctccgcctc ccgggttcaa gcaattctcc tgcctcagcc tcccgagtag ctggaattac

cattacctg gitgicctit ccgigccact ggaittiggi cicattaig thagcigaa ctggagacca igtggaggaa ccgaggaccg gaaggicca citigggicca iticcagicco aggeaaitat gitatagcia cigagggat cicigitgic attaaacgco acagtgicci egrecette ettettagga agececteaa tettececae caaceteetg agaggaggee tetaacaaae aeteetttae agacagttt gggttgcac gctacttatg agaatctaat gtctgatgat ctgtcactgt ctcccatcac occcagatgg gaccatctag ttgtagga scaageteag ggeteceact gattetaeat tatggtgagt tgrataatta tittattata ttaataeatt atggeegggt geagtggete iggggaatgg agtgaacctg ggaggcggag ctcacagtga gccaagatcg caccactgca ttccagcctg ggcaacagas aggaggoog aggoaagaga atcacttgaa cocaggaggt ggaagttgoa gagagcogag atcgoatcac tacactgoag gogotggat occgggcacc ccagtgatgt gcotggagag ggcccaggot totgaggcga occacacott gocagtcagg cagitoticae tgaaggecce eccagtgect ggeectgige tgggigiggg gatacaggga geeggggaga ceaggagag aggotgaggo aggagaatgg ogtgaaocog ggaggoggag ottgoagtga gtogagatog ogooactgoa ctocagoot ggetetett actigecagg ggtagaeteg gecectiggaa caagetecag aggeacagee caggagetgg tggaggeoa Agetcagca ccacggacte gaaaaacace etggocaact gcoggotgga aaaaacaace aegaeggtgg ccoeggeet acceteage etgtgaaact gacaaateac etgegaagge tecaetgggg caggttetgg gggagggtg ggaatetgee egggacaag gggtgagggatgcattcca agcaaaggag acaaaacctg cataggagtg aaatagtccc gtgtgtttgg ggacciticag ggaciticcae tggaggeagi tigeaaggaa gaagagage atgtgfaggi cagggeagaa gttaggicag ocagoacit tegebagace agglegegeg ateacaaggt caggagatte agaccatoot ggotaacatg gtgaaacoo geccagging cgcatgaggc actgcatect cteategece acetgggeag agaagggeat gatgteettg aaageaatge gagaagaga ctaatgootg taatoocago aotttggag googaagoag gaggatoaot tgatgooagg agttogaga igácccacat ggicccagaa gcaagggcot ggggccticc tgggtticcg tcctggtggt ttcagcccat caggaaggtc. gggaaaccc cgtctctact aaaaatacaa aaaaattagc cgggcatggt ggcgcgcgcc tgtagtccca gctacacggg goodgegca taggottott caaacgoott caggocaggg acagooott totggatggo cacgoocago accatoocaa icaccigiaa icccagcaca tigggaggcc gaggiggig gatcacgagg tiaggagitt gcgaccagcc iggctaacac catectgaa accaacecet cacacacet teettagaaa aattgtettt cacgaaacea mmmmmm mmmmmm acgoctacg gagtgtotte cetgettece tggecaggat gggtagagtg gtggeagetg gaeceetggg geoceetee gregaga gaaatgcacc ttgrggatet getecaaaag etgaaagaaa egegtateat gaageeaca eagacageae stototacto aaaatacaaa aaaaattago tgggogtggt ggogggogo ogtagtocoa ootaottggg aggotgaggo ggocaccge atacacagee eggtatgegt tgtaggcaga acteatggag aaggettga gettgggeat egtgtgtgc gocciggai citatcactt gacacticca agacacagtg ggtgagagaa ggcaaggaic agagagaaag aictgictaa agatoccotg accagtggco tggttotoca gtgcotgcac coctagotgo ocatagtogt cactgotgoc aaccagagag atocaggico accogaacti otgoagoago agoaccatgg totocacoig giactigica tiggggaigg tgogoaggaa ggectgtta ggaaccgggc cteacaggag gaggtgagea getggtgagt aagegaaget teatetgtat ttaeagetge cagcotggcc aacatggtga aaccccatct ctactaaaaa tacaaaaatt aggcogggtg oggtggctca ogcotgtaat ccigggigac agágccagac igictcagaa aacagaacaa aacagaacac gagacteta teteaaaaaa aaaaaaaga aaagaaaaat tateeaggea tggtggtggg tgeetgtaat eesagetaet getecotgi acactgigic agcateacee ecaggetetà ggitgeceat aagceagita catggigagi agceacatee itgaaagcti ggcattotot gcagagoiga tigotgotgo accaggagoc ottgiggcaa ggootagggg cottotigto sgaggatac tgccgcttca cgctgagcgt ctcgctgctg gccgcatagc taatctatgg gaggtcccgt tcagccatt ecectectg teagateage ageageatea gattettgta ggagettgaa ecetaetgtg aaetgeacat gegagggate iggicatita gagtgaccgg agagtgocca cicigcical cicaggatig gcigiticico cigacaggag gtgciggggt

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LECOAFMAHT MPKLKAFSMS SAYNAYRAVY AVAHGLHOLL GCASELCSRG JAGATVVVVF SSRQLARVFF ESVVLTNLTG KVWVASEAWA LSRHITGVPG RVYPWQLLEQ IHKVHFLLHK DTVAFNDNRD PLSSYNIIAW DWNGPKWTFT **SSSDDYGQLG VQALENQALV RGICIAFKDI MPFSAQVGDE RMQCLMRHLA** ORIGMVLGV AIOKRAVPGL KAFEEAYARA DKEAPRPCHK GSWCSSNOLC ASCSENEHGY HLFQAMRLGV EFINNSTALL PNITLGYOLY DVCSDSANVY /HISYAASSE TLSVKROYPS FLRTIPNDKY QVETMVLLLQ KFGWTWISLV ATLRVLSLPG QHHIELQGDL LHYSPTVLAV IGPDSTNRAA TTAALLSPFI sagaaggtot cottggagot ctatgtggtg ttgocot

caccaccact ctcagctaac ttttgtattt ttagtagaga tggggtttcg ccatactggc caggctggtc tcgaactcct ggoctcaaga ticaagaaa tictootgoo toagcotoot gagtagotgg gattacaggt gootgooaco acgootggot aaittitgoa tititagoag icititicit icigagacag agicitigcic tgicgcocag gatggagtigc ggtggcgtga tctiggcica cigcaaccoc tgocicotgg stagagggcc tegaagaggg agaggaatga gggcaaccac aggccaggca ggaacccatg gggaaggatc cataagccaa aatcgggctg agggtcaaig agggcaggga gaggccagca ggaaactccc aigggaaggg gcagggagtc agtgctcagg cccatgagag tggaggcagg gatctggaag cagctctgga aagagaaggaa ggctggggca ggaaccacgc tgggcaggg cototgagec aggaggaag aaggaaagge aggeaggaga gactgggatg atgtggagca gtotatgggg tgggaageaa iggggaggag aggagggga agcotgctcc ggggaatcac ctacottitc agaggaagtg gggcaaaagg agagaagag caggggctc agatcagagg ggaggggact gagaatggga ggttaaacca cgagcccaca gcctgcctgg gaactggaaa ggaggggttg tggtccaagg tacagggcaa gaataagcac agagacagga ctgacatcag caaggtgagg catgtcagca gcaagigaa agccaggigg gggcaggggg cigaggggg cataaattcc aaggaaagac tctcatagga ggactggtca eccetgacit gigaciaaag agcagigace acceaagaga iccaggggge aggeageeit gggggggaca geageicitg gotggtgtga attocagotg tggotgtggo agtggaaaag gaggocagaa aggatgaaag gtggggagoa gggcaagga gaacctctgg agggaggagg gaagtggagg gcagcagggg tacagctgag tggcagtagt tcccaaggag aatgggtttt cceacatgoc ccagoccaga ottgootgaa gggagatggg caaaggtotg aggotocago ttaccatggg caccaggaaa gggeteagea gggeggetgt ggtggeagea eggttggtge tgteaggeee aateaetgee ageaeegtag gggaatagtg gagcagcagt gggcaggact ccagggtgat ggccactccc tcactaccct ccaccagagg attggggcta atacaggaag gaictgocca gootoccaa gggattacag goatgagoca cagogocogt coaggatgte cattoctaac aaaggoaacg ការបាលបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការបាលការប ataaagaagg actgcaaagt aggatttgga tacctagaag gtgccccagc tcacagcgaa agcaagagtg gtggggaca unimminimo ceactéctet aagceacage gagteectaa ggateteege agagaagtge tatettegga ettgeattit iccactgcac tecagectga gtgtcagagt gagactgtgt etcaaaaaaa aaaaaaaa aaaatcacaa gtcacctaag gggattaca ggogtgagcc cccgcgcccg gtgcccggcc gggacttgca tttcatgagc gtatctctga cttcagtgag gaatgagtta gaagaaattt aagactaaaa tcagggggaa gccttaggac actgatggga gaatctagct gaggggtgat raggagitca aggecagtet aggeaacata gigagacete tatetetaca aaaaatacaa aaattageca ggeatggigg acatgootg tggacccago tacttaggag tatgaggtgg gaggattgot tgagootggg agacagtgag acaacattgo aaaatgteac aaagggeaeg gtgeeteatg eetgtaatet eaeeaetttg ggaggeeaag geaggtggat tgettgagee igggigtoot ittitggggg gaggaiggag gggacaaggt atcactotgt cacccaggot ggaatgcagt ggtgcaatot agacagggtt teaceaegtt ggecaggetg gttteeaaet eetgaeetea tgagetgeee aeettageet eecaaagtge aaaagagget tttgttgtgt agggaggtaa ggteaatetg ggeettgetg ggteeatgat gtggeaatgt tgggeeagea cageteactg caacetecae eteccagatt ceageaatte teetgtetea geeteccaag tagetgggat tacaggeaca

LR92

Coupled Receptor G Protein-GPCRB3

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VLGSSTWSPV QLNINETKIQ WHGKNHQVPK SVCSSDCLEG HQRVVTGFHH CCFECVPCGA GTFLNKSELY RCQPCGTEEW APEGSQTCFP RTVVFLALRE HTSWVLLAAN TLLLLLLGT AGLFAWHLDT PVVRSAGGRL CFLMLGSLAA GSGSLYGFFG EPTRPACLLR QALFALGFTI FLSCLTVRSF QLIIFKFST KVPTFYHAWV QNHGAGLFVM ISSAAQLLIC LTWLVVWTPL PAREYQRFPH LVMLECTETN SLGFILAFLY NGILSISAFA CSYLGKDLPE NYNEAKCVTF SLLFNFVSWI AFFTTASVYD GKYLPAANMM AGLSSLSSGF GGYFPLKCYV ILCRPDLNST EHFQASIQDY TRRCGST	gagcaacatg atctittiga agtactigac ggtgtcgttc ttgacggtca cgaagcacag agtgttgatc atgctgttgc tcatggcgat A gcactogacg atgtagaag cagtgaggta gtgcttctoc ttcacaaaca cggtggggaa gaagtcgcgc acgatggga agcactogacg atgtagaag gagtctctc ttcacaaaca cggtggggaa gaagtcgcgc acgatgggaa agcctttcct gcggcagcgc agcctttgagaa gggtcttgggaa gggtcttggggaa cgatggacgg catagcacag ggtcattggt accacggggc catagcacag ataaagaga agtagaccag gagatcttgg catagcacag ggtcattggt accacggggc cacagaattc tatgccaaag ataaagaga agtaggactt gtagtagagac tgctggtcca caggccagat ctggccagaa tggccagaa tagaacagaa accagatcttt catggcttt gacaatgacg aggaccgtc cggtgggaa gaagacttcag aggatgggcaa tcagggagaa accagagaa accaaggcaa tcaggccagt ggctgtttgg cacttcattc gtggtctcag cggatgggaca atagccagat acctagggca agaacacaag tggaggcagc c	MGFMDDNATN TSTSFLSVLN PHGAHATSFP FNFSYSDYDM PLDEDEDVTN SRTFFAAKIV IGMALVGDML VCGIGNFIFI AALVRYKKLR NLTNLLIANL AISDFLVAIV CCPFEMDYYV VRQLSWEHGH VLCTSVNYLR TVSLYVSTNA LLAIAIDRYL AIVHPLRPRM KCQTATGLIA LVWTVSILIA IPSAYFTTET VLVIVKSQEK IFCGQIWPVD QQLYYKSYFL FIFGIEFVGP VVTMTLCYAR ISRELWFKAV PGFQTEQIRK RLRCRRKTVL VLMCILTAYV LCWAPFYGFT IVRDFFPTVF VKEKHYLTAF YTVECIAMSN SMINTLCFVT VKNDTVKYFK KIMLLHWKAS YNGGKSSADL DLKTIGMPAT EEVDCIRLK	ggacgagge gccggcgcc atgtggagct gcagctggtt caacggcaca gggctggtgg aggagctgcc tgcctgccag gacctgcagc tggggctgtc actgttgtcg ctgctgggcc tggtggtggg cgtgccagtg ggcctgtgct acaacgccct gctggtgctg gccaacctac acagcaaggc cagcatgacc atgccggacg tgtactttgt caacatggca gtggcaggcc
	AX147788	LR114	BC014241
	WO0034334- hFB41A	194904 WO0034334- hFB41A	G Protein- Coupled Receptor MGC7035
	194904	194904	194905
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CTGAATGCGC GCGGCAGCG GGCGACGCGC CCTTGCGCAG CCTGGAGCAA

GCCAACCGCA CCCGCTTTCC CTTCTTCTC GACGTCAAGG GCGACCACCG

ICCGGACTAG TTCTAGACCG CTGCGGGCCG CCAGGCGCCG GGAATGTCCC

LD22826

Coupled Receptor

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G Protein-

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MGC7035

ATCAAGGAAG AGGCTCACCG TAAGCCTGGC CTACTCGGAG ACCCACCAGA GCTGGTGCTG GCCGCGGTGG AGACAACCGT GCTGGTGCTC ATCTTTGCAG TGTCGCTGCT GGGCAACGTG TGCGCCCTGG TGCTGGTGGC GCGCCGACGA ZTGAGCGGCA GCGTCACCAT CCTCACGCTG GCCGCGGTCA GCCTGGAGGG CATGGTGRGC ATCGRGCACC TGGAGCGCGG CGTGCGGGGT CCTCCGCGGC SGGCGCGGGC AGTGCTGCTG GCSCTCATCT GGGCCTATTC GGCGGTCGCC CGCCGACCAG GAAATTTCGA TTTGCACACT GATTTGGCCC AGCATTCCTC GGACTGGTCA TTGTGATCAG TTACTCCAAA ATTTTACAGA TCACAAAGGC CTCTACAACA TGACACTGTG CAGGAATGAG TGGAAGAAAA TTTTTTGCTG CTTCTGGTTC CCAGAAAAGG GAGCCATTTT AACAGACACA TCTGTCAAAA ICACACCTGG CGAGCTGTGG CATGCTTTTA AACAGAGTTC ATTTCCAGTA ATCCACAGCG TCGGTAAATT AAGGGGTGAT CACCAAGTITT CATAATATTT SOCTOCATOA GTGCACCOTG CITTAAGAAA ATGAACCTAT GCAAATAGAC SECCECECE CEACTECCTE CCTEGTACTC AACCTCTTCT ECECEGACCT SCTCTTCATC AGGGCTATCC CTCTGGTGCT GGCCGTGCGC TGGACTGAGG CATECTIGATE CAGAACTIFCA AGEAAGACET GGICATETIGG CEGICECTET SCTCTGCCTC TGTGCGTCTT CTTTCGAGTC GTCCCGCAAC GGCTCCCCGG CCCTTTATA AAAGGATTTG TTGGCCAGGT GCAGTGGTTC ATGCCTGTAA CCTCCCTGCT GGGCCCCGTT GCCTGCCACC TGCTCTTCTA CGTGATGACC GAGAGATCTC GTGGGATGTC TCTTTTGTTA CTTTGAACTT CTTGGTGCCA COGCGTGTC CCAGCAGGAC TTCCGGCTCT TCCGCACCCT CTTCCTCCTC ICTICTGGGT GGTCCCCTTC ACATTTGCTA ATTCAGCCCT AAACCCCATC ATGGTCTCCT TCTTCATCAT GTGGAGCCCC ATCATCATCA CCATCCTCCT GAAATGACTT GTCGATTATT TCTGGCTAAT TTTCTTTATA GCCGAGTTTC

Homo sapiens

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TCCCAGCAGT TTGGGCTGAG GTGGGTGGAT CACCTGAGGT CAGGAGTTCG AGACCAACCT GACCAACATG GTGAGACCCC CGTCTCTACT AAAAATAAAA	GGAGGCTCAA CCACGAGAT CTCTTGAACC TGGGAGGCAG AGGTTGCAGT	GAGCCGAGAT CGTGCCATTG CACTCCAACC AGGCCAACAA GAGTGAAACT CCATCTTAAA AAAAAAAAA AAAGATTTGT TATGGGTTCC TTTTAAATGT	GAACTITITIT AGTGTGTTTG TATATGATCA AATTTAATAA ATATTTATTT ATGACTGTTC AGCAAAAAAA AAAAAAAAA AGGGGGG	MSPECARAAG DAPLRSLEQA NRTRFPFFSD VKGDHRLVLA AVETTVLVLI FAVSLI GNVC ALVI VÅRBRP PGÅTACT VI N I FCANTI FIS AND VIT AVDVI	TEAWLLGPVA CHLLFYVMTL SGSVTILTLA AVSLDRMVCI VMLQRGVRCP GRRARAVLLA LIWGYSAVAA I PI CVFFRVV PORI PGADOF ISICTI IMPT	IPGEISWDVS FVTLNFLVPG LVIVISYSKI LQTTKASRKR LTVSLAYSRS HQIRVSQQDF	KLEKTELLEM VSFFIMWSFI ID ILLILLIQ NFKQDLVIWP SLPPWVVAPT FANSALNPIL YNMTLCRNEW KKIFCCTWFP EKGAL TDTS VKRNDLSIIS G	ITYSAISDEL RDKVRFPALL RTTPSADHHV EAMVQLMLHF RWNWIIVLVS	SDJ YGKDNGQ LLGERVARRD ICIAFQETLP TLQPNQNMTS EERQRLVTIV DKLQQSTARV VVVFSPDLTL YHFFNEVLRO NFTGAVWJAS ESWAIDPVI H	NLTELGHLGT FLGITIQSVP IPGFSEFREW GPQAGPPPLS RTSQSYTCNQ	ECDNCLNATL SFNTILRLSG ERVVYSVYSA VYAVAHALHS LLGCDKSTCT KRVVYPWQLL EEIWKVNFTL LDHOIFFDPO GDVALHLEIV OWOWDRSONP	FQSVASYYPL QRQLKNIKTS LHTVNNTIPM SMCSKRCQSG QKKKPVGIHV	CCFECIDCLP GTFLINHTECP NNEWSYQSET SCFKRQLVFL EWHEAPTIAV	ALLAALGELS ILAILVIFWR HFQTPIVRSA GGPMCFLMLT LLLVAYMVVP VYVGPPKVST CLCROALFPL CFTICISCIA VRSFOIVCAF KMASRFPRAV	SYWVRYQGPY VSMAFITVLK MVIVVIGMLA RPOSHPRTDP DDPKITIVSC	NPNYRNSLLF NTSLDLLLSV VGFSFAYMGK ELPTNYNEAK FITLSMIFYF	TSSVSLCTFM SAYSGVLVTI VDLLVTVLNL LAISLGYFGP KCYMILFYPE
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RNTPAYFNSM IQGYTMRRD
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clcgocggga tcccgggtga ttctgacat agtgtttggc tttggggctg tgcgggctg tgttggaaac ctcctggtga tgattcaat
cctccatttc aagcagotgc actotcogac caatittct gtggctctct tggcdggcg tgtttcttg gtgggtgga tgattcaat
cctccatttc aagcagotgc actotcogac caatittct gtggctctct tggcdggcg tgattcttg gtgggtgga ctgtgatgc
cttcagcatg gtcaggacgg tggagagctg ctggtattt gtggatggt tttgacttt ccacacctgc tgtgatgtgg catittgta
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gaatttgctg ttggtgtgct tattataact cagccatgaa tcctttgatt tatgctttat tttacccatg gtttaggaaa
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STCGAFYIPS VLLII HEGHSHSAGS PLFFN SLVLPICRDS CWIHP		togoggite teege oggetaatti titga tgaaccccog acete ogaacctca eteao		agcatcatct greet agcatcatct acatt ctgagtgtgg acatg gacaggtact gggc gccgcgctga tgatc ttctggagaa gccac		attagatgcc gagac cctcatgagt ggatc ggagagtttg taagt ttgtttgagg attgt aattcaaat aaaca EKMLICMTLV VITTI IXIVMDRWKL GYFLK IMILTVWATS IFTSN LYYRIYHAAK SIYQK
NTSQISYTIY SSLCSLNSSL IICWLPFFVV VPFRKAS	gagagaagca ctggagccag agagtctcag	cctcccgggt ccaccatgcc atgctggtct gaattacagg	aatggaacac ttcgaaggcta ctgaaacaag acccaagacc acccaagacc	gcctgccaac catgccctg tgaggtgtgg cattgcctg ggccaagagg	daottctcq qaattctqata ttgtaaaactt gtttgaaaag aggagaacgt gattctgggt tctgagcatc	taaaaagctc ttttccagag acttggttca ttgttctgtt tgtgatacat SMAIRPKTIT VAVLVMPLSI ARKRTAKRAA FYIPLTLILI
AQEEMSDCLV TAHLITGSAG KILGIILGAF EEFRQAFQKI	atcgaatgtt agtgagactt accgtccaca	gcaacctccg caggcactca catgttggcc	accaacagaa gtgagaaacc acagtgtaga tggctataag	agenceacea cagtgetegt acttectetg acetetgtgt ggaagagge teatetecat	atatococtt tttaccagaa cttttgcaag ctaccacaga tagatcaccc tcctggggct tgattgtgggg	
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131		NP_000857.1	MDFLNSSDQN	LTSEELLNRM	PSKILVSLTL	SGLALMTTI	NSLVIAAIIV	TRKLHHPANY P	Ношо
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			STIYSTEGAE	YIPLALILL	YYKIYRAAKT	LYHKRQASRI	AKEEVNGQVL	LESGEKSTKS	
			VSTSYVLEKS	LSDPSTDFDK	IHSTVRSLRS	EFKHEKSWRR	OKISGTRERK	AATTIGLILG	
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Receptor 5-HT2C

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NTKELNSPIL FAFSHLESSD GEAGRDPPAK DVMPGPRQEL LCAFWKSDSD RGGHWATEVC	QVLGSKNGST	PIQGSRTTIH	GLELYFLWR	WSFLGPVTFI	VEGLEIFDDR	NHULUSLLSL

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EMR1 Hormone NM 001974

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			PKOKFERPIC	VSWSTDVKGG	RWTSFGCVIL	EASETYTICS	CNOMANLAVI	MASGELTMDF	
			SLYIISHVGI	IISLVCLVLA	IATFLLCRSI	RNHNTYLHLH	LCVCLLLAKT	LFLAGIHKTD	
			NKTGCALIAG	FLHYLFLACF	FWMLVEAVIL	FLMVRNLKVV	NYFSSRNIKM	LHICAFGYGL	
			PMLVVVISAS	VQPQGYGMHN	RCWLNTETGE	IWSFLGPVCT	VIVINSLLLT	WILWILRORL	
			SSVNAEVSTL	KDTRLLTFKA	FAQLFILGCS	WVLGIFQIGP	VAGVMAYLFT	IINSLOGAFI	
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	GPR30		gcgagtgaaa	attcaaatgg	ccagtagggg	gcgcactcgg	aagtggccgc	cccgcatgag	
			gcagttcagc		gtccggggag	ggaggtttat	tctccgcctg	cacgagactg	
			tgaaatccgc	aaccatgagc	aggagagcg	gccctggtgg	ggaagaggcc	accaacatct	
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Receptor 2

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Receptor D1

Dopamine

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			aaaacccgga		gaccatgago	cgtaggaagc	tctcccagca	gaaggagaag		
			aaagccactc		cattgttctc	ggcgtgttca	tcatctgctg	getgeeette		
			ttcatcacac	acatectgaa	catacactgt	gactgcaaca	tacagaatgt	cctgtacago		

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	Homo sapiens	Homo sapiens
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	NP_000787.1	MM_000797
	Dopamine Receptor D3	Dopamine Receptor D4
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tgtcaccgcc tgcaccccgt tccggccccc agacgcccct

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	LAVTKNKKLR SVVGSIENIV YDPRTYTCIF AEVRNFLTMF AVIYGLINEN RAHACPAVEE SKRAASGHIKP SSNPKPITGH DNPELSASHC	agaggaggag ttgttggcga gggagcctgc ccaccatggt tccccatgg ccagaaa	togagoccat gattoccoat gattoccocca acatactcccca acatactcccca gaggattocc tcctgaatga acaaaatcta acaaaatcta gatgacacaa gatgacacaa gatgacacaa gatgaaga tcattgaaga tcattgaaga acatgacacaa gatgaaga acatgaaga ccttgaaga ccttgaaga ccttgaaga tcattgaaga
	VDLIGNSMVI CQMVGFITGL LPNMYIGTIE PAGONPDNQL FIAYFNSCLN HARDQAREQD STHHKSVFSH KPDSVHFKPA TTSHPKPAAA ADLPDPTVVT		atccagaggg ctgcccaaca ctgcaacaaa agggatcaacc acttgtaca cttggggaga cttggggaga gccattctg gaactccgag gggggaatca gaactccgag gggggaatca ttccagtgcc ggcatgaga cttcagtgcc ggcatgaga ctccagtgcc ggcatgaga ccagtgcc ggcatgaga ccagtgcc ggcatgaga ccagtgcc ggcatgaga ccagtgcc ggcattaga aaactgaga ccagtgcc ggcattaga gccatttaga aaactgaga ccagtgcc ggcattaga aaactgaga ccagtgcc ggcattaga gccatttaga aaactgaga ccagtgcc ggcattaga aaactgaga ccagtgcc ggcattaga aaactgaga ccagtgcc ggcattaga aaactgaga gccatttaga aaactgaga gccatttaga aagtcctcat
	MECAMVITIV VIGGWDLSQLQ CO ITWINTVLAAV LIWITKVLAARD ELINWLYLAAY EARTLARARA HESSSAYRKSA SVHFKGDSVHF KETADXPKPA TAASQLESDTI AAASQLESDTI A		
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		agagcaaaat	aacaggaata	tatttaggaa	agattccctt	gctgctctgc	tggaccccct	
		gaaaggtgca	tgcacttgaa	gctttgagat	tccagtgcca	ccgcaaccca	tcttcatcca	
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sapiens Homo

Glutamate Receptor

LGSEIRDSCW HSSVALEQSI HTEGNYGESG DDYFLKLRLD EKGDAPGRYD GQIKVIRKGE IESIIAIAFS KPTTTSCYLO AFKTRNVPAN FTPKMYIII KSVSWSEPGG TKTLYNVEEE KGLPPPLOOC LOAASKLTPD VONLLQLFDI SILISVOLTI PPOHLOMLPI GPGSSSVAIQ KLOSPEVRSF RYNWTYVSAV LPKARVVVCF EENYVQDSKM GVSGEEVWFD RSVCSEPCLK IPVRYLEWSN GYVCPFTLIA MSAWAQVIIA GLLIMSCTYY LSVTVALGCM AGAGNANSNG GKSLTFSDTS PLFLAEPALP GLRSLYPPP EDELEEEED ADPVLLPNIT QARAMLDIVK PNADLTGCEP KKICTRKPRF KILTTCEAUS TFLNIFRRKK KPLTKSYQGS LPPHLTAEET YEHEREGNTE DRLLRKLRER EVEANGGITI KRICTGNESL LDFLIKSSFI KIOMNKSGVV CYILLAGIFL LGVVAPLGYN VLAGPGGPGN RTKKPIAGVI AMFHTLDKIN ERFKLLQEYV PGHLLENPNF RIARILAGSK EVYLICNISN GDGKLPCRSN RVPSAATTPP LPDGQSLPPG FLRVVPSDTL IYSNAGEKSF ADRDEVIEGY AMKPIDGSKL TCKACDLGWW FSTAIPDFHA HEGVINIDDY PVVKSSSREL FVPIYFGSNY ETACNOTAVI REQYGIQRVE PGSPSMVVHR MDQLQGVVSN SPPADDDDDS EGLCIAHSDK EFSLIGSDGW LCPGHVGLCD KENEYVQDEF CYSALVTKTN MYTTCIIWLA TISDVVRMHV HRLSVHVKTN RDEKDGINRC DLSDKTLYKY FWOHRFOCRI RYDYVHVGTW TLIFVLYRDI MPILSYPSIK SAMRRIGVVG TNTRNPWFPE AHGLONMHHA IMILOYTEAN RLLVGLSSAM EDAQPIRESP QOPPPQQKSL QLSTFGEELV MDAFKELAAQ VSCCWICTAC WTLIIMEPP KPERNVRSAF GOVPKGOHMW CLGILVTLFV FNEAKYIAFT EFIRDSLISI POIAYSATSI

gtggctgagg ctgttcccag cgtggcatco cacctgctgc catgcgctgg tcacgccaca actggtgtta ctatttcaga ggcgagacag ctdctdcaga gagetgette tggggggcc cggaacaaca aagatcatgt cgctatgact gagattctcc SSTL cctcttgagg ggccatggct acaggagtee SVILRDYKQS gtggggtgct gctgggtggg caatgagcac ccgtgacccg caaggacaca tgctgatgga cactdccatc tgacaagtcc gggcgactat ctgtgtggcc ggtgcgagcc ggatgcccgg cagtgatggt tactatcacc cctggaccct gagacttggt gtggtcctgt acagttgctc tcagccgtgg aggtggccaa ccaagctgag tccaagccaa aggcctctga cccgcaacat cctgggtggc ctgctgaggg LCTPPNVSYA tgctgccgct accgcatcaa gtgatgctcc ttgagggtgt scottctga tgcggccagt VPSSPVSESV ctcctggcac tttgcactgg cgtgcctcac qtctccatcc gtgtccactg accctggagg gcagaggact cacatcctcq gcgacccatg tctaccagtg cctgacttct gaggetegtg gtcctgttca gccagcttca agtgagggg ctccgggctg aagtttgatg cgcgcggcct tttgcctcct gaattctggg tagctacgcc ctttgagcta ggtggcaggc catcagtgac tgcagtgtac cacccddctc gctcaacgtc FRDSVASGSS ggccatgctt cctgggtgca ctacagtgat cacagtgcct tgccatgagc agcccactct gctgcttgcg gaaggtgctg gggcggccca ggactttgtg ctggacctat gegeeteaat cggctcttat ccgcgtggct DSPALTPPSP ccatgggatc gcccagccaa tgcaccagaa agegeetgga tctgccccga ttggcggttc tcccacagat actttqccq gcttcttcaa gcattgaggc aagtgggccg ctgccagcca tggagagtgt cctcctaccc gccggaaccc gagactgcgc ttgtggtcaa gcccaacac aggactttgt ctggcgtgcg agcaggcact ageceagtge

Metabotropic NM 000839 Glutamate

Receptor 2

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	Metabotropic NP_000831.1 Glutamate Receptor 3	Metabotropic NM_0 Glutamate Receptor 4
	3095	9600

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		tctgccgtct	gtcttgcccg	cctgcccgcc	tgcccctcct	gccgaccaca	cggagttcag	
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3096	Metabotropic NP_000832.1	MPGKRGLGWW	WARLPICLLL	SLYGPWMPSS	LGKPKGHPHM	NSIRIDGDIT	LGGLFPVHGR P	Ното
	Glutamate	GSEGKPCGEL	KKEKGIHRLE	AMLFALDRIN	NDPDLLPNIT	LGARILDTCS	RDTHALEQSL	sapiens
	Receptor 4	TEVQALIEKD	GTEVRCGSGG	PPIITKPERV	VGVIGASGSS	VSIMVANILR	LFKIPQISYA	
		STAPDLSDNS	RYDFFSRVVP	SDTYQAQAMV	DIVRALKWNY	VSTVASEGSY	GESGVEAFIQ	
		KSREDGGVCI	AQSVKIPREP	KAGEFDKIIR	RLLETSNARA	VIIFANEDDI	RRVLEAARRA	
		NOTGHFFWMG	SDSWGSKIAP	VLHLEEVAEG	AVTILPKRMS	VRGFDRYFSS	RTLDNNRRNI	
		WEAEFWEDNF	HCKLSRHALK	KGSHVKKCTN	RERIGODSAY	EQEGKVQFVI	DAVYAMGHAL	
		HAMHRDLCPG	RVGLCPRMDP	VDGTQLLKYI	RNVNFSGIAG	NPVTFNENGD	APGRYDIYQY	-
	•	QLRNDSAEYK		LRIERMHWPG	SGQQLPRSIC	SLPCQPGERK	KTVKGMPCCW	
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		DPSHSVVDFQ	DORTLDPREA	RGVLKCDISD	TSTICTTGAS	MLLMVTCTVY	AIKTRGVPET	-
		FNEAKPIGET	MYTTCIVWLA	FIPIFFGTSQ	SADKLYIQTT	TLTVSVSLSA	SVSLGMLYMP	-
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	tgc	gcg	cgg	င်ရှင	g	g Ca	gag	cag	aca a	atg	cca	aga	tgt	aaa	ctt	aag	act	tct	ttt	tgt	SVL	RVE	RCV	FMR	IXS	ADR	EGF	AMK	DNG	T CK	Σ	RIA	EVZ	FVP	909	KEN	LYD	SIO	AIE	PNS
	tccagttgc	ggo	စ်သ	Ü	tgt	att	gedededdad	gcc	ctcatgaca	agaaacatg	acagttcca	ıtga	ttcccctgt	attcaccaaa	ttt	ccttgaaaag	ataacacact	atttcctct	ctacttattt	ttcaccatgt	TIT	REQYGIQRVE	EEEEGLVRCV	DKTLFKYFMR	CIAHSYKIYS	LLGSDGWADR	HRFQCRLEGF	GYAGLCDAMK	YINVGSWDNG	EYVEDEYTCK	FIIYRDTPVV	ALVTKTNRIA	HDYPSIREVY	TCIIWLAFVP	TVVRMHVGDG	LSIHINKKEN	PDAGPKALYD	SOGSIMEQIS	AEIOPLPAIE	VDSGSTTPNS
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Homo sapiens

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Metabotropic NM_000843 Glutamate

Receptor 6

3098

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Homo sapiens

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Receptor 7	-	TEVOALIOKD	TSDVRCTNGE	PPVFVKPEKV	VGVIGASGSS	VSIMVANILR	LFQIPQISYA	
		STAPELSDDR	RYDFFSRVVP	PDSFQAQAMV	DIVKALGWNY	VSTLASEGSY	GEKGVESFTO	
		ISKEAGGLCI	AQSVRIPQER	KDRTIDFDRI	IKQLLDTPNS	RAVVI FANDE	DIKOILAAAK	
		RADQVGHFLW	VGSDSWGSKI	NPLHQHEDIA	EGAITIOPKR	ATVEGFDAYF	TSRTLENNRR	
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Ното	sapiens

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Metabotropic NM_000845 Receptor 8 Glutamate

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gagaccatca tggtgccagt ctaggaccc attctcctat ttatcagtcc accetctaga aacagaaagc aatttttagg cagctatggt caaattgaga

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	Homo sapiens				Ношо	sapiens		:																
gtg gcacttgggc scct ctgcaacaga igaa aaaagtggaa	SNVL VMISFKVNSQ P LALD YVASNASVMN	LCWQ YLVGKRTVPL LADL QGSDSVTKAE	ATGP SANWAKAEQL /KAE TEKSDYDTPN		SNSK LP acca aagtgaccag A	ygtg agtgggaggg aacc tggatagacg			ccag ccctgggcca stcc ctggcgtatg		tttc tccgacgcct	-	tgig ttegecagea toat eeettgaaac		aggo egtactetet		catt gttggaatta tgag cagctaaagg				gaga gcatttcgct			agca acgccaagag
tgtgtcccag tcaccctgtg aaccccatct gctatgccct ctctgccgat ggaaaaagaa	> 0	IGLAWLISFI LWAPAILCWQ ILYCRIYRET EKRTKDLADL	WSSSRRSTST TGKPSQATGP ESPGEEFSAE ETEETFVKAE		LCRWKKKKVE EKLYWOGNSK atctgaagac cccggcacca	tettgggetg ecegtgggtg eteteceage accadaaace			ctreceege gecereega gacgeatege getetagtee				tctttcctat cacagetgig gotatatoge tattattoat		aaaccaaagt catgccaggc		tgggtattac atacaccatt octotosa gtatoatgag		caatctatca acaactaaat					ccagtcggaa gaaaagagca
	yaacaycaay cracc PLERHRLWEV ITIAA IFSMNLYTTY ILMGR	KRTPKRAGIM IGLAW AFYIPVSVMT ILYCR	LAQRERNQAS WSSSR LQVVYKSQGK ESPGE	_	TFRKTFKMLL LCRWK ttccagtctt atctg	aggagteteg tettg occatoocca etete			ctgcctgtgg cttcc cagccgtcct ggcgo				ttccagaact tcttt qcqqtqqaca qqtat		ctttattcca aaacc		rrgercarca rgggr Craddadata cetot		attctcactg caatc	_	aataaaagat ttcga			accaccaggr ccagt
	TVNGTPVNHQ PLE SLACADLIIG IFSI	SITRPLIYRA KRT PTITFGTAIA AFY	RSCLRCPRPT LAQ DEDKPATDPV LQV		NPICYALCNR TER atctttcagc ttc	agaacttcag agg cadaccooto oco			cgcgctggga ctg ccadttcdtg cad		-		ctactgccgc ttc gacgccatt gcg		ccctcagtgt ctt		crgrttccca trg		tatttacttc att					cgatgcagac acc
atggtcctgg tattggttgt accttcagga		LLVISFDRYF DECQIQFLSE	KRKPAHRALF TTCSSYPSSE	YLLSPAAAHR NPSHOMTKRK	YWLCYVNSTV ctattgcagt	gaggcagaga tecooogeto	ggggtggagg	ccacgggggc	acctcaccaa	gtgtggtggt	acaagcgcat	ccatggccgc	totactocat	ccagactgtc	tacttgcctt	gctttgtgca	ctctggtgta	CCaaaagaaa	tgccctatca	acatccagca	ccatcatcta	ggtgtccttt	atccaaaccg	ttgaccccaa
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	Muscarinic Acetylcholin	e Receptor M5			Tachykinin	Receptor 3																		
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				GHSMKQEMAM	NMVTNSVLLN	RMTSLKSNAK			-	
	TSYLLSSSAV			PFALYLLSES	VLSFGNSCVN	GHMIVTLVAR				
	FNYNEIDPSL	PNHILYMYRS 1	FVGCFIFCWF	RKRLAKIVLV	NEHTKKOMET	KSAHNLPGEY				
	YYYHIAKTLI	FLIPLAIISI '	IHSVLIFLVY	YPOTDELHPK	DNSSFTACIP	FSEVARISSI				
	SVLLAVPEAV		MQTSGALLRT	RYRAIVNEMD	VETLTALSAD	VIQLISVGVS	٠.			
sapiens			LLLICVPVD	FISNLAAGDL	NSAMRSVPNI	NIMLVKIFIT		Receptor	·. ·	
Homo	LLIITVGLLG P	VIRCVIPSLY 1	PASDGTTTEL	VPEGWERDFL	VTTGANESGS	L MPSKSLSNLS	B NP_002502:1	Neuromedin 1	3380	199
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	gattttggcc	atgt	gaagcaggaa	ggcacagcat	ttactaaatg	caattctgtt				
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	tgaccaccgg		gtctctttcc	tgccctctaa	aaggagatca	ctctgctgga				
sapiens	ctccagcgga			tcagtcctca	tcgtgggcgt	taaacctaaa	٠	Receptor		
Homo	ggacatcgat A	cgagagggag (cttgcagggg	ggacagtaaa	gcttgcccgc	gtgctgtgag	B NM_002511	Neuromedin I	3380	198
		VDEYS		RRNSKSASAT	PRDPSFNGCS	TRSSRKKRAT				
	VVFDPNDADT	ESMI		_	FRWCPFIKVS	KRFRAGFKRA				
	YNPIIYCCLN		WKYIQQVYLA	LTAIYQQLNR	CWLPYHIYFI	MIIVVMTEAI				
	LKAKRKVVKM		GITLWGGEIP	LIMGITYTIV	VIILVYCFPL	PKOHFTYHII				
•	TLCEVQWPEG	MPGR	AFLLAFPQCL .	KIVIGSIWIL	LKPRLSATAT	VDRYMAIIDP				
	ASIYSMTAIA	ITAVE	WYFGANYCRE	VNFIYALHSE	DASMAAFNTL	NYFLVNLAFS				
sapiens	LAHKRMRTVT	IIWII	AYGVVVAVAV	PSWRIALWSL	WANLTNOEVO	PVASPAPSQP	:	Receptor 3		٠.
Homo	LSSSPSALGL P	WLQLLDQAGN I	GAATGAVETG 1	AVNLTASLAA	IDGGGGVGAD	٠.	NP_001050.1	Tachykinin	3378	197
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Neuropeptide NM 000910 Y Receptor

Type 2

3404

ctggaattca gcattatgag

gtggatctaa

tgtgaaata gtagtaggtt

gcagagcctg

agttggttgg

tttacttaac

aacaaaatgg

aagataaggc

tttcccattt aactggctgg

tgaaaactga attcctggaa

ggctcacaag

ctgctgttta

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> ggattgagga gaatttctc

tcaaagcatt

attggtatta

gttaggacct ccactgaaca ttgttcattc cgaatggctt

ttcctggagt

tegetgetee

acgtttggtg

aaccaattgc aattacagga

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aactgaaatt

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aatdcaaacc tagataacaa

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cgctttatgg

cctgttgtta atctaatctt ttagaaggaa

gagagactgg cactaatcca

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tgatacttt

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aaattccaag

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ctttttgaac

gctctgctga ctctgggcat

Ношо	sapiens
Д	
LAYCSI	EWKMGP

SEVSVIFKAK

RCEQREDAIH

NPLLYGWMNS NYRKAFLSAF

HIIAMCSTFA

KEYKLIFTVF KNLEVRKNSG

PNDSFTEATN

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> Neuropeptide NP_000901.1 Y Receptor

Type

3404

ctttattgct tctgcagaag

tttactatct

cagaattgcc

ctgattcata tagtttgtct

tcatggccat

attctgccct

agttcagtat

tataagctgt

atgtgttgag

gtggaacttc aagaaacatt

ggacactagg

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ctttctgata tcacagtctt tactgtaagt catacaagtg

gaaaatgaga

gccataaatg

tcaggtgaaa ctctctggca

ccatccaaaa agagtgggcc

aactcttcat

ttaacagcaa accatggcta

tgtcaggtat catatgataa aacatcccat atctaataat

ttttgccatc tgttctcccc ttccagtgtt tggttcagca ttgctgagca gcaggtattt

202	3405	Neuropeptide NM 005972	atgaacacct ctcacctcct ggccttgctg	stoccaaaat ctccacaagg tgaaaacaga A	Ното
		Y Receptor	agcaaaccc tgggcaccc atacaacttc	: tctgaacatt gccaggattc cgtggacgtg	sapiens
		Type 4	atggtettea tegteactte etacageatt	; gagactgtcg tggggggtcct gggtaacctc	
			tgcctgatgt gtgtgactgt gaggcagaag	gagaaagcca acgtgaccaa	
			ccttctctga	tgcctcctct gccagccgct	
			tacaccatca tggactactg gatctttgga	gagacetet geaagatgte	-
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			gagaatgtct tccacaagaa ccactccaag	y gototggagt toctggcaga taaggtggtc	
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			ttocagtact gecteceact gggetteate	s ctggtctgtt atgcacgcat ctaccggcgc	
			ctgcagaggc aggggcgcgt gtttcacaag	y ggcacctaca gcttgcgagc tgggcacatg	•
			aagcaggtca atgtggtgct ggtggtgatg	gtggtggcct ttgccgtgct	
			tcaacagcct	caccatgagg ccatccccat	
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			atctatggct ttctcaacac caacttcaag	y aaggagatca aggccctggt gctgacttgc	
			cagcagagcg ccccctgga ggagtcggag	y catctgcccc tgtccacagt acatacggaa	*
			gtctccaaag ggtccctgag gctaagtggc	s aggtecaate ceatttaa	
203	3405	Neuropeptide NP_005963.1			Ното
	-	Y Receptor	CIMCVTVROK EKANVTNILI ANLAFSDFIM	M CLLCOPLTAV YTIMDYWIFG ETLCKMSAFI	sapiens
		Type 4	QCMSVTVSIL SLVLVALERH QLIINPTGWK	K PSISQAYLGI VLIWVIACVI SLPFLANSIL	
			ENVEHKNHSK ALEFLADKVV CTESWPLAHH	H RIIYTTELLL FOYCLPLGFI LVCYARIYRR	
			LOROGRVFHK GTYSLRAGHM KOVNVVLVVM	M VVAFAVIWLP LHVFNSLEDW HHEAIPICHG	
			NLIFLVCHLL AMASTCVNPF IYGFLNTNFK	K KEIKALVLIC QOSAPLEESE HLPLSTVHTE	
			VSKGSLRLSG RSNPI	•	
204	3406	Neuropeptide NM_006174	gaaaggctat cggtaacaac tgacctgcca	a caaagttaga agaaaggatt gattcaagaa A	Ношо
		Y Receptor	agactataat atggatttag agctcgacga	gtattataac	sapiens
•		Type 5	taatactgct gccactcgga attctgattt	t cccagtctgg gatgactata aaagcagtgt	
			agatgactta cagtattttc tgattgggct	t ctatacattt gtaagtette ttggetttat	
			ggggaatcta cttattttaa tggctctcat	t gaaaaagcgt aatcagaaga ctacggtaaa	
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			tctgtcttgc	g gatgtttggc aaagtcatgt gccatattat	
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ttcatcaaaa	gaagacct	agtcagctct	cctgaagaaa	aagagatctc	tggatgccac	aggcatttca	aatccaattc	cactgtcttc	KTLATENNTA	NOKTTVNFLI	ILISIAIVRY	LLSSRYLCVE	ENEMINLTLH	ILPENFGSVR	TILILVEAVS	GIKADLVSLI	cccgcgcagc	ggagatcgga	cacgggttct	ccggagcccg	gtcttcgcca	actcctgccc	ccatgcgcct	адсадасаса	gcaacgcgtc	tctactccaa	gcaacacggt	cggtgcatta	tgcccgtgga	gctgccgcgg	ccagcctgag	tgtcccgaag	cggtgcctat	gaggactggt	acaccttcat	ccaacaagct	ggggcgagca
 gagttattca	accedereca	ctctgtaaga	tgagataaa	aagaataaaa	tgctgttagt	tatttcaaat	ctgttgtctt	gtcccttata	MDLELDEYYN	LILMALMKKR	QCVSVLVSTL	VELQETFGSA	GLSNKENRLE	ERPSOENHSR	KRSRSVFYRL	NPILYGFLNN	tcaagctcgc	cgcgcggttt	cccgaggaac	agcccggagc	gagacagata	agacdcdcc	ccagcgccca	gaccccttcc	aacgcttcgg	aacaccgaca	ggcacggtgg	ctgcagagca	ctgctggcca	ggcgacgccg	ctcaacgtgg	aagaccctca	gccctgctga	cagcacgccg	atacaggtca	accatcatcg	tgcacggtcg
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									Neuropeptide NP 00	Y Receptor	Type 5				٠		Neurotensin	Receptor	Type 1										·		<i>\$</i>	. •					
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Homo sapiens	Homo sapiens					
pa totogtatoa ctagottgog gocaggtoat gatgtggcoc oggaagotggg coatgagtgg toggtcatgg agtcoggado coctgagocg gococtggtgg cotcacagot caaacgocacactoc caccatctgc aggtggtgaacc gtgtatotet caataaaggt ggcogaaggg cotcgatgtg g sElDVNTDI P TALEVVGTVG NTVTAFTLAR KKSLOSIQST VHYHIGSLAL SDLITLILAM NV HHPWAREGDAG CRGYYFLRDA CTYATALNVA SLSVERYLAL CHPFKAKTLM IS AIWLASALLT VPMLFTMGEQ NRSADGQHAG GLVCTPTIHT ATVKVVIQVN NV VISVLNTILA NKLTVMYRQA AEQGQVCTVG GEHSTFSMAI EPGRVQALRH TA AFVVCWLPYH VRRIMFCYIS DEQWTPFLYD FYHYFWWTN ALFYVSSTIN NV FRHIFLATLA CLOPWWRRR KRPAFSRKAD SVSSNHTLSS NATRETLY	accigicate gactgecage eggetgaggg egggggtete aggaggttge agaagtaceg tacagagtgg attigeaggg teceedace ofteragag attatetace geaceacet	tgagccccaa cacagicty etgeccccgc atctgctgct tcctgcccct cgggctcaq gtcaccatcy tggggctcta gggctctt gtcatgtacy tcatcctcag ccaccatat ttacatctt aacctggccc tggcccag	cettecaggg cacggacate etectggget tetggecgtt cagteattge cattgactae tacaacatgt teaccagcae gtgtggateg etatgtagee atetgecaee ceatecgtge gcaaageeca ggetgteaat gtggecatet gggeeetgge	tigecateat gggeteggea caggitegagg etacecetea ggattactgg ggeceggigt tegtececgt getegteate tetgtetget teegeetget etegggetee egagagaagg tgetggtggt agtggetgt ttegtggget	cccaagggct gggggttcag ccgagcagcg agactgccgt cggccctggg ctacgtcaac agctgcctca accccatcct acttcaaggc ctgcttccgc aagttctgct gtgcatctgc tgtctgaccg ggtgcgcagc attgccaagg acgtggccct cggtgccct acttccaagg acgtggccct	oggicatotac gectaacaca gageteacac aggicaetge gggccatotac gectaacaca gageteacac aggicaetge gggccetgag cateagagac etgggatggg etttrecetg cagagagaga acetagigac ateatgggac aggicaaagc eccagagacaga ctaaagcgc ectectggtg cagggccgag
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NP_002522.1	NM_000913					
Neurotensin Receptor Type 1	Opiate Receptor- Like 1	(OPRL1)				
3408 8	3452					

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	Homo sapiens	Homo	
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tgtgcagccgg cagaccccga aggcctcatc cagcgagagg accagccct atgggcagct ggtgggagaa ctttgcttga gaagctggtg agatggctct ccagcatgag gctgtgagga	KVTIVGLYLA ILLGFWPFGN NVAIWALASV ISVCYSLMIR QPSSETAVAI SIAKDVALAC	aacacagccc cacgcagctc cgggctccgc cgggctcccc cgaccttctc ttttgttgac ggggagtggat tcacatcatg ctaccttcc catgtacctg gactgcagtg gactgcagtg gattgcgagtc catgtacctg catgtacctg catgtacctg catgtacctg catgtacctg catgtacctg catgtacctg catgaggggct ttgaggggttct cctgggtttt tcaagtgggc tcaagtgggttt tcaagtgggc tcaagtgggttt tcaagtgggt	
cgactccacc tccctggctg tgcacggtgc ttcaggagac tggaccgtca gcgtgaccac gctctgtttg acagcctctc tgtggaagga acaagcctca gacagcctca	SHGAFLPLGL LLTLPFQGTD VRTSSKAQAV LFSFIVPVLV VFVLAQGLGV RDVQVSDRVR	cgcgtccgcg gggacgcagc tgggcagcagc ccgctgcctg gattcccaaa ctttctgcgt tgttttgcta tcctgctgta ccatgctcta acaagacagt acaagacagt acaagacagt ttatttgttg cagagaggaa ttatttgttg caggatgcag caggatgcag caggatgcag gctcggctgcag acaagacagt acaagatgcag acaagatgcag acaagatgcag cagga	
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cctggaggac gtccaggtgg tctgaaggtt gggccaacc gtgcaatgaa tgtctcagga tcgtttcct atcctccaa gctgtgttgc tggggacgcc gccaacggg tgcttgagcc	NLSLLSPNHS KMKTATNIYI LTAMSVDRYV LVEIPTPQDY ITRLVLVVVA FLDENFKACF	gggtcctggc gaccttctgc ggtctgccac cacggaaatt cagtgcctgc atcggcagga gctctgtggg gctctgtggg atcggaccc acaggaaccc acaggaaccc acaaggcatt atcattctt atcatatcat atcatatcat atcatatcat atcatatcat atcatatcat atcatatcat atcatatcat atcatatcat atcatatcat atcatacat accatagacat atcatacat atcatacat accatagacat accatagacat accatagacat atcatatct atcatatct atcatatct atcatatct atcatatct atcatacat accatagacata accatagacatagacata accatagacatagacata accatagacatagacata accatagacatagacata accatagacatagacata accatagacatagacata accatagacatagacata accatagacatagacatagacatagacata accataga	
ggagctgcca aggagaagt ggaccgcacc gcttgactct ccctccagcg gtggggcagg agtggaggcc cagtggccgt agtcctgctc tgagcttgct tggcagggct gttg	EVIYGSHIQG LVMYVILRHT YYNMFTSTFT AQVEDEEIEC SREKDRNLRR NSCLNPILLYA	caggocogago gcoctagg tccagocogo gcottotgat atatgaacca agctgttgta tgatcogag tggccaccct gtgagocogo tggttctcgt ttaaaggaag tcagaactgo tcagaactgc tcagaactgc tcccttttt tcagaactgc tcccttttt tcagaactgc tcccttttt tcagaactgc tcagaactgc tcccttttt cagaactgc ctgcaaggaac	
coctgagott gggccacccc gctgactgca cctgactgcat gcttctcagt ctgttcacaa gattctctgg agccagaggt gccacagaggt gccacagaggc gggtggggcc	MEPLFPAPFW VCVGGLLGNC ALCKTVIAID VGVPVAIMGS RLRGVRLLSG LRFCTALGYV KTSETVPRPA	atgacccagg atggcctccc gtgctgagct ttggcgctgg gcgacgtccc ggctgctcgg atgtggatcc gcttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatctgg gcgttatcatga ttgaaacctg ttgaaacctg ttgaaacctg ttgaaacctg gcccagggat cagtctccca cagtctccca gggaaactt gggaaactt gggaaactt gggaaactt gggaaactt gggaaactt gggaaactt gggaaactt gggaaactt gggaaactt gggaaactt gggaaactt gggaaactt	
	NP_000904.1	MM_000273	•
	Opiate Receptor- Like 1 (OPRL1)	Ocular Albinism 1 (Nettleship- Falls) (OAI)	
	3452	3513	

gtaatttctc taatactgac

tccatgcttt tttgtaacat caaagaaaac atacccatca ctttctattc tctattaata aaaaattaat acatacaatt

attcaattct attatattaa

aaatagtaaa aataccagaa

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	gotocagaco coatattoot cotocoactooc	100000	agagereer		gttttctgag gctggctgta	taaaatagtt atgactg	_	I LRAMAACDIL GCLGMVIRST	: EWWLFCYAVD AYLVIRBAG	I AIPHYVTMYL PLLLVLVANP	1 LVLIICWLSN IINESLLFYL	PARTICISTRE OSPRKEIQWE	1 LSEGSDASTI EIHTASESCN			taatggaaaa	s actgaaaaga gacctcatat				tatctcaaga	s cttggtgact caggccttgg				ctcaccaacc		tttcttttgt	c aagtcaagtc ggaattccac	s gtgtttgtgt tttttgtctg	agtcagaccg	a ttcactctgc tactatctgc	a tgccagccgt ttagggaaat	t gacctagaca tttccagaat	tgagttccta	aacagaaatc	a attragttca ataaaattca	
•	tatgaagggg atgtgctggg ctttagaact gtgttctcac			ttcatgcaca cacgtgtgag	gccttagttg ccactaggaa	tccttgggga agtagttaaa	TPEPRPRIOP MASPRIGIFC	GRRPAGPGSP ATSPPASVRI	WPAAFCVGSA MWIQLLYSAC	EGAAMLYYPS VSRCERGLDH	YTENERRMGA VIKIRFFKIM	IWFIMGILNP AQGFLLSLAF	PASGKVSQVG GQTSDEALSM		accttggage ctacaatgag	aggcctagac gcaggatctt	tcactgggca aaacaccttc	aggeetetge etteagaagt	aatcctgctc tcagaacctc	tottcattgo gggaatccta	gctctaagag tttcatcatc	tgacttttcc tttcaagatc	ttgtgtgcag ggtctctgcc	ttgggctcat cagctttgac	-	ttgctgttcc aaatattatt	_	_	aaatctttaa gtcccacctt	gccgcaacat attcagcatc	ccagaatccc ctacacaaag	tcttgcggta tatgaaagaa	ctattattta tttctttcta	ttccattaaa agctcagaat	ttgaaagcac agatactttg		atctctagca ctgccatcca	
	catggagace t			•	agctgctcta g	aaggtccaca t	MTOAGRRGPG	LALGLIQLLP G	SVSDMNHTEI W	AWGLATLLCV E	ASLLKGROGI Y	LKPVRTAAKT T	HPSPLMPHEN P	HGDL	gaacagtgtt a	tcacagatga a		-	cctccagatg a	tactgtatgg t	tacgtgccca g				•		-	•			taccatattg c			aaattgcaca t	aatacaacac t		cataaatatc a	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
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							Ocular	Albinism 1	(Nettleship-	Falls) (OA1)					UDP-glucose	Receptor	(KIAA0001)																	•				
				•			3513								3544							•	-															

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	Homo sapiens	Homo sapiens	
aggcacagtt gatttgaaga gtatttcatg ttttttctga ttacgtcatt agagaaacta aataagatga aatgggaaag tttacattaa gaaacagac aaaactaaat ttctttcaaa	PSSKSFIIYL P FFGLISFDRY KCIELKSELG SSRNIFSIVF DPIIYFFLCQ	gtetgegegg A ceaggeacag tegeeteetg tecagtgaga gggeegggag teaactttag tggeacgetg teagaggagg geecetacae aaggeeggg	accccgcgg gctcctggcg gcactcgcgc gtttcaggtg gctgtccatg ccgcaccgac gcaggtgcac cttcatccag
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	atactg DESCSONILI SITFPFKILG FIQSVSYSKL FVALEWIVEL IARIPTKSQ		• •
tctagtatgt tgatgaaggg agcactgcaaa tagcactttg taatgagcct taatgagcct aatattggca agaccattt tgagtgcaaa ggattttact tctttctctg	aaatgtttta MINSTSTQPP KNIVIADFVM YKIVKPLWTS RKWHKASNYI VEFVCFVPYH	tittaagget ctggeetege cgeegetege gaceteaget cgeacgegte gttggeetetgg gtggaeceag ectecgaeae	gccaacgca cgcaacgaga ctgtagcaga ctcttcttct ctgccaggt ccctggaac gcctggaac atcttctctc
	NP_055694.1	MM_000916	
	UDP-glucose Receptor (KIAA0001)	Oxytocin Receptor	

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	NP_000907.1	NM_002564
	Oxytocin Receptor	Purinergic Receptor P2Y, G- protein coupled, 2 (P2RY2)
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		NP_002554.1	MM_005767	
		Purinergic Receptor P2Y1	Purinergic Receptor P2Y5	
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3597	Purinergic Receptor	NP_004145.1	MEWDNGTGQA TAVYTINIAL	LGLPPTTCVY ADLLYACSLP	RENFKOLLLP LLIYNYAOGD	PVYSAVLAAG HWPFGDFACR	LPLNICVITO LVRF1,FYANI	ICTSRRALTR P	Homo Sapiens
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			AVVVAAAFAI	SFLPFHITKT	AYLAVRSTPG	VPCTVLEAFA	AAYKGTRPFA	SANSVLDPIL	
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	NP_005039.1	NM_000316
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YKIAALVEYS YYAKDEWPEG ACVGVWIMTL FIMIGCYLVI ENSYNPWGAF RSLSNINSEM						
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NP_005283.1	NM_006143				NP_006134.1	NM_016602
G Protein- Coupled Receptor GPR18	G Protein- Coupled Receptor	GPR19			G Protein- Coupled Receptor GPR19	G Protein- Coupled Receptor GPR2/CCR10
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		Receptor		_	LTCICVDRYL	AIVRPEAPAA (CROPACARAV	CAFVWLAAGA	
	-	GPR20		VILSVLGVTG SRPCCRVFAL	TVLEFLLPLL	VISVETGRIM	CALSRPGLLH	QGRQRRVRAM	
				QLLLTVLIIF LVCFTPFHAR	QVAVALWPDM	PHHISLWYH	VAVTLSSLNS	CMDPIVYCEV	
				TSGFQATVRG LFGQHGEREP	SSGDVVSMHR	SSKGSGRHHI	LSAGPHALTO	ALANGPEA	
260	3858	G Protein-	NM_005294		taatcagage	agccaccett	tttgcctctt	ggcatttggc A	Ното
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		GPR21		_	tatccagact	atggcatatg	ctgacctttt	tgttggggtg	
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				gcctgtatca gcattgatag	atacattgcc	attactaaac	ctttaaccta	taatactctg	
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261	3858	G Protein-	NP_005285.1		YLETVNFCLL	EVLIIVFLTV	LIISGNIIVI	EVEHCAPLLN P	Ношо
		Coupled		_	SCWVPSLSLL	HHPLPVEESL	TCQIFGEVVS	VLKSVSMASL	sapiens
٠		Receptor		•	VTPWRLRLCI	FLIWLYSTLV	FLPSFFHWGK	PGYHGDVFQW	
		GPR21		_	APAALIVCET	YFNIFRICOO	HTKDISEROA	RFSSOSGETG	
					FYILWLPYII	YFLLESSTGH	SNRFASFLTT	WLAISNSFCN	
				CVIYSLSNSV FORGLKRLSG	AMCTSCASOT	TANDPYTVRS	KGPLNGCHI		
262	3829	G Protein-	NM_005295	atgtgttttt ctcccattct	ggaaatcaac	atgcagtctg	aatctaacat	tacagtgcga A	Ношо
		Coupled			caccaatatg	taccaaccac	tatcatatcc	gttaagcttt	sapiens
		Receptor		_	tcttatgtta	gaaattgtgt	tgggacttgg	cagcaacctc	:
		GPR22		Τ.	catgaaatcc	aacttaatca	actctgtcag	taacattatt	
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gaaaacaaga cacttttatg tacaccaaaa tacttcaggc aagaagaaag caagaaagaa atgtcacaaa gcagtggtgg ataattgcc tccggcgagc gtcttcagga tgtctttatt gttttaaata ccaccatttt tgtttttag tcatggctta agacaaaaat ttcaaaaggt gaagctgatc ccctgcctaa	gtcacagact ag YQPLSYPLSF QVSLTGFLML ICVGCIPLTI VILLISILESN ILTMGRAVML MISIWIFSFF YHLLVQIPIF FFTVVVMLIT MSQSSGGRNV VFGVRTSVSV VLNTTILCLG PSDLLVKLRL EADPLPNNAV IHNSWIDPKR	agagaattca aacagtgaag gtaggattca ttgctccttc tgtcccagg atcatcgcct tgtcccagg atcatcgcct tgtcccagg atcatcaac tctcggtagt atggatgcca atagtcagt tactggtact tgcccacc ttggtgatct gccatcggt tactggtact tcccttca gccagactca tgcccctcc tggtgatct tcccttca gacactgacc tctactggt gacactgacc tctactggt gacactgacc tctactggt gtctccagc cagacacca gtggtcatca tccctttgtgt gtctcggtct tctttgtgtg atcaccagc cagacacca tgtctggtct tctttgtgtg atcaccagc cagacaccac tggtccagac gcagacaccac tggtccagac gcagacaccac acgacaccaaccac ttggtcctgt cggtgaagcc acgacaccaaccac
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	G Protein-	Coupled	Receptor	SLC/MCH1		•		G Protein-	Coupled	Receptor	GPR25		-					-	37					•			G Protein-	Conpled	Receptor	GPR25			G Protein-	Conpled	Receptor	GPR3		• ,	
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				FIFFAFITAS AYVIMIRMIR	VVKOIIFIPA LNITICHDVL PE SSAMDENSEK KRKRAIKLIV TV	PEQLLVGDMF TVI AMYI I CF	
				YFLIKSQGQS HVYALYIVAL	PFVYYFVSHD	FRDHAKNALL	
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7	4052	Proteinase-	NM_004101	cggcacagga gagcaaactt	ctacagacag accaaggett co	ccatttgctg A	Homo
٠		Activated		gaactgaggt gaaattgtgc	tccatgattt tacagatttc at	ataacgttta	sapiens
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20	4052	Proteinase-	NP_004092.1	LLLLLPTFCQ SGMENDTNNL	RGAPPNSFEE	FPFSALEGWT P	Ното
-		Activated December 3		PEESASHLHV KNATMGYLTS	YLLVEVVGVP	ANAVTLWMLF	sapiens
		Receptor 3	•	FYTNLAIADF LFCVTLPFKI	FGEVLCRATT	VIFYGNMYCS	
				LELLACISIN KYLAIVHPET YKGLPKHTYA LV		LKOEYYLVOP	
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	Ното	sapiens	•																																	Homo	sapiens		
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Homo sapiens

gtg A Homo

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DPIMYFFVAE

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LSAKSEL

CGKRLKGPPP SFEGKTNESS

ctcqtcttca atcttcatga tatatcatga aacccactgg gacaagaat acctgggaca cccaaggcca tgatatggag atgttgtgaa agtgaacatt gacagtcaca ggacggtgaa tcgtgtggaa tacatgttcg gcagagaagg gtgccctacg caccttccc attettgett acacagtago aagggagaac ggccctaact gggcccacag tggtccttgg ttccgcttcg cacataggct gtgtctatg tacccacagt aagaagctgo tgacctcttc ctteggteec ccctgtcatc ctgcggcaag ccatccccta ctcctcactc teceaegtte gaacacgagg agaatggggc aatgaatggg acttggctaa taaaatggaa aggtgtgtgt attctagtta acctcctgat tgtaggcagg tgggggagg cgtccagcac catggcgctg ctatgggcag cacacagaag gatctgctgg gagccaggtg tgaacgaagt cacccctagt tggcacagaa ctacatgttt cttcgtcttc catgagcaac cctgcagtgc ttttgtcatc cttcgcagca cccttcgag aattgccctg taggcgtctc tgtctagcac tcttctccta agaccaaaag tatgattatc ggggttgggc ctatggagag ctgtgcagaa attaatgagg tgatctggag catcttcaq ggaatggagg agcaactcat aaacaacac cctggggtct gageteagge cadccatgaa tgctggccgc tctacgtcac tagccgtggc tgtgtaagcc tcacctgggt tegettteet agggctccaa ccatctacaa ccaccatctq agacggagac agctgtacag tggtacgcag tgcatggata tgggcggtga acaacgagtc agtcagccac tccccgaggg tcttttctq tcgcagcagc gctactgaga agttaattac ccaatgtggc ttcacccacc accgtgtcca tggccgacta ttaagaaata tagggataag aatagcaaga tgagattggg tgctctagca tgcatgctca ggagcagcdc caagacctac cctcccaact gatgcaggaa ggcctcactt accagggctg ttecteaege ctgctcaacc tacacctctc tccaggtaca ccggaggtca atgattatca cagcagcagg atcatggtca aagagcgccg ctgagtggct acaagggcca gcgacgggtg tttgccaccc tacgtggtgg ggcgttgcct caqttctcca gcactttgta taatgtaact tcaaggccag catggtcatc attctacatc gttccggaac gcattcagat ccaatgaggg actgatatt gttctttgcc ggcctctgct taggactctg ggacatccac tgcccctcct tacatacata ccgacacgca atgcagtcat agctggagcc cgggtcagcc cttctccaat tgagccatgg cccatcaac caactacatc cagcacctc ggagggcttc catcgagcgg tgccatcatg caccaactaa cacgeteaag caccatcccc ggccgctgcc catcccacca tttttttt tgtcccagct atgctggatg tctcagaccc ttgagattgg tctggaaag tgcttaataa cctggtcctg ggccaagttc tggtccactt ccagcgtggc ccatcccagc aagacctgcc cagccacagc atatctatcc cagttgttt taacatcaat tagctaggca ggaatgcagg aggtcccgtg actacctggc gtggcttcac gatgcaattt Eggteetgge Lgaacaagca gtgacgatga ccttaatttt gcctgagaag gcgggatgtg gtgtgtttca agagtcatcc ggagcagcca tctacgtgcc tgctgggctt gcacgcctct gggagaacca caccccact tegactacta ccgtcaagga ggtcacccg

Rhodopsin NM_000539

4254 Rhod

Homo sapiens	Homo sapiens	Homo Rapiens
tccttgggga agcagttgct tgaggtgtca agaagctcta tcagctccta gatagattga atgagcagag gaatgggaaa GFPINFLTLY P NLEGFFATLG	KEAAAQQQES PAFFAKSAAI agtgcaagctc ccggaagctgc atcagcctga tcggacgct agcagtcag tggaactcag tggaactcag ttcgcccttc gactactcca ttcgcccttc gactactcca	aaactgcaga tatgccctgg agggagaagg tgttccagga ccagtggccc gcacagaag aggcctcagg gttttgttac cctaataata tcaccttct tttagctccc
gg aaattccact gggcctacct ct tgccagacaa gcccatcttc tc caaaaagctg gccacatctc cc ttctccatat aagcaaagcc gg caaattgggc cattaaaagc tg ctttcacact ctatccacag cc tgggatggct ggattgagca ga ggtggaggag gcagtcctgg YY IAEPWQFSML AAYMFILIIVL GG FTSTLYTSIH GYFVFGFTGC GE NHAIMGVAFT WWALACAAP	HFTIPMIIIF VAEYIFTHOG DEASATVSKT gagtgaggat ctgtgggggat tcttctctt tggctcttgc tccggcgctg cagcgttggc gcacccgtag cttctgcctt tggggacatg tcaccatgc tcaccatgc tcaccatgc	ag acytgactic catcicococo go ccacgatcaa tgccatcaac go agtgcctctc accgcagaag ag tgagcccag gccaggaggc cc cagacactca ccaccttcc gg attcagaag acaccaggct gc ccctacact caaggctgag aa atgtaagggg gtacagtgag aa atgtaagggg gtacagtga cc ctcacccc tcccaccttg cc atgtccacgt gtacacatta ac tttca .3G ISINTITIES FCKTPELRTP
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tactcgaaga tgttcatggg agtccattct gaattaagct gctttaccca tgttggtatt aactgccagc ccaagcagca aacccca VTVQHKKLRT GEIALWSLVV	EGLOCSCGID ATTOKAEKEV YNPVIYIMMN O2921 agagacagct ccactggctt ggactccctg atgccctcgt gccaggctca ccatcgctc tgggttgggg agggggacag ccctctct tgggttgggg agggggacag ccctctctct tgggttgggg	tygtyccoge gcaatgagat accgaaccaa gtcctgccca gtggatcct agccagatgg aaagtcattc atggatagat tacgttgtac gagtctccaa acttacaagt
Rhodopsin NP_000	Retinal G NM_002 Protein- Coupled Receptor RPE	Retinal G NP_002
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CCTLDYSKGD LLGWGPYAIL SPQKREKDRT	agctcccgag cgggcaccat	a Laga Laga	gcacggagca cttctgtgcc	gcagaaatgg	cta	acctgctgaa tggtcgcct	at go	stot	ccarggrgcr tctaccttca	gtgg	ttto	atcc	cacc	gaat	gtto	ttca	gcatcatctg	aaggetggg	300a	agaa	C	LFR	LGILCAFREL	LFQYCIMANY	VGCW	MOON	
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FWAALPLLGW QKLGKSGHLQ NAINYALGNE	ggaccctgcg gggcgccctc	yctactactg atgtgacgtg	agagcagaca caacataagc	attecteegg	ctggtcagaa	caacgagaag ctcctcccto	ctgcactcgc	caacttcatc	cayyycyyyc ctggctgctg	aagaaagtac	tttgtgggct	cgcarccarc	aaatgaagtc	tggcatccac	caatggggag	cccactgcac	ccagggcacc	ayarraayay gararrattat	gggatgtgag	ggaagagaag	ttc AAHSTGALPR	PGRMVEVECP	KLKVMYTVGY	SSDDVTYCDE	EGWGSPALEV MRKLRTOFFR	GLVVAVLYCE	
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SQLAWNSAVS SFFNFAMPLF SISPKLOMVP	a cgaggcegg gcacgggcag	actggagee	rgccrgcagg qqttqtqaqq	gtggaggtgg	cgaaactgca	gttaatgtga atgtacaccg	tgtgctttcc	ttcatccttc	tgcatcatgg	gccatctcct	totocagoca	tecatectga	agaacccaag	ctcctgctga gctatggaga	gacgtactat	caatggcacc	gccagccact	geagggeeart	aggccttgga	ttggttcgtt	aaatggtgcc MRPHLSPPLO	QPVPGCEGMW	LACGVNVNDS	LEVSFILRAL	HILLAISEES PVIISILINE	SPEDAMEIQL	NSTKASHLEQ
		actggageee	tgcctgcago qqttqtqaqo	gtggaggtg	cgaaactgc	gttaatgtg	tgtgctttc	ttcatcctt	tgcatcatg	gccatctcc	totocagoos	tccatcctga	agaacccaa	gctatoda	geogreete	caatggca	gccagccac	gragggrea	aggeettgg	ttggttcgt	1.1	QPVPGCEGM	LACGVNVND	LEVSFILR	HTLLAISE TATESTA	SPEDAMEI	NSTKASHL
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RPE	NM_002980	a chagagaco	rgccrgcago qqttqtqaqo	gtggaggtg	cgaaactgg	gttaatgtg	tgtgctttc	ttoatcott	geaceae	gccatctcc	totocagoos	tocatoctga	адаасссаа	cretgeto	googtocto	Caatggca	gocagocao	สวาร์นนูลาน	aggeettgg	ttggttcgt	NP 002971.1	I	LACGVNVND	LEVSEILR	HITTELLATER PALISITIA	SPEDAMEI	NSTKASHL
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II Type 2		VNIVVVTLFC	COKGPKKVSS	IYIFNLAVAD	LLLLATIPLW	ATYYSYRYDW	LFGPVMCKVF	sapiens
Receptor		GSFLTLNMEA	SIFFITCMSV	DRYQSVIYPF	LSQRRNPWQA	SYIVPLVWCM	ACLSSIPFFY	
		FRDVRTIEYL	GVNACIMAFP	PEKYAQWSAG	IALMKNILGE	IIPLIFIATC	YFGIRKHLLK	
		TNSYGKNRIT	RDOVLKMAAA	VVLAFIIWCL	PEHVLTELDA	LAWMGVINSC	EVIAVIDLAL	
		PFAILLGETN	SCVNPFLYCE	VGNREQOKLR	SVFRVPITWL	QGKRESMSCR	KSSSLREMET	
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ic Receptor		agtgaggtgg	agctggactg	ttggtttgat	gaggatttca	agttcatcct	gctgcctgtg	sapiens
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Receptor		IFRLRPWDAT	ATYMFHLALS	DILYVLSLPT	LIYYYAAHNH	WPFGTEICKF	VRFLFYWNLY	sapiens	
		CSVLFLTCIS	VHRYLGICHP	LRALRWGRPR	LAGLICLAVW	LVVAGCLVPN	LFEVTTSNKG		
		TTVLCHDTTR	PEEFDHYVHF	SSAVMGLLFG	VPCLVTLVCY		LPGSAQSSSR		
		LRSLRTIAVV	LTVEAVCEVP	FHITRTIYYL	ARLLEADCRV	LNIVNVYKV	TRPLASANSC		
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LLLLGRRARA AAGADAGPGP EPCATLVQGK FFGYFSAAAV gtatgcagaa ctagacccag tcccctccaq TRTYLGVESF VEYLVVGNRN NEWSSWSACS NIOMMTREHL AKAORGLPGE LSIHKLPASG DVPSSSAPPQ CTLVAAFLHF caccaagaac catcatcgac gcgggccagg gggcggacgg aaaagaatta RGDVCLRDAV APGVEGGGCE OTGDPAAEEW WGSCSVTCGA WKETPAGEVA LYRNIGSFLA RAGASIWSSC RDKAPKSSFV ccaggacctg ggacagcaag agcccacggg ggagettege cagtgctggg acacccccat AWDEWSPWSI NEVOILSNLI FTKAKGYSTN gaggaggat accagaggo tgagctcctc TYQFDSFLES tgctggggcc ggcacagggc ggtacccgcc ttcttcaata GPPGPTDDFS GGPAAGPLAP LOTRTRICLP ELOOFGEPAP NNSAVCPVHG CPGRAVDGNW VDGKWQAWAS YYSPTPGDVQ DAYOVTDNLV EASVEVVGTV NOTCILWDET GOTOTRNKVM PALVVAI SVG. DGITDKKLKE DFPNHSLTLK aagacatgtt. tgggccagga taggcccctc caggactgag CDEDNEGAVI VMVHCILRRE IAACRTATIT cggggcccag ggaagtcgcg gcctccggaa cgtcgccgct acagggcccg gcgggcagat cqttttttaa MEKATLPSVT ctgggccacg ctcgcgggca VPCSGPGRVR QFLQMRRQQP PQHDGLRPRA TGGWKLWSLW GECTRDCGGG tecttttett TPCACLGGEA DARRREELGD KOTKFCNIAL VIGERMKDLR ANVSKLHLHG SPRYPGGPLP ctggagcggc gacaaggagg ccgctgcagc cggccacgca gcggccaggc ggcctggcac ccagggccc SGPLREQRIC TRDCFLOQCP RNMTEIFRRA ILVFNKLVSK LACRSVLNKD ccctgggaga atcccgctgg ccggccagcg ccgaggccca YIRCVSIDYR VESTGLTEAD ILAQLSADAN aagcggcacc cacagacacg agcgtcccag ccccagggg TORCPEPHEI EFAHMYNGTT IISSNALILI KRFLCLGWGL atttttctc RSSHPCGIMO GAECQGHWVE SVILINECLS AVVLVNMVIG LMTDFEKDVD TLYMKVAKAP SSRSQSLRST FGGNPCEGPE GPODEYROCG EGIAYWEPPT GDLLSTIDVL LFRLVEDEVD PEDRVTVSKS CLCDRLSTFA GHLRNRLIR tgtgagctcc agcggagag aggaggcggc cagogogoc acctcatgg CVSSSYSTOC gcacaccgg caacaagagg ggagctggag gggtgggcgg cgcagacgg cgctcagacg cctcctcggg tggcccggcc cccacacct PRSLRTPLEI gggcgccacg cacccatcca tgcagcacgc VWILAPLLLL LLYAFVGPAA LTQDRGGHGA GGSFONGHAO GPPTNENSLP gggagaggtc ggcctcaggg gggggaatct TLRNPDPRRY DEVLRLCDPS APLAFLQASK GEGWOTRTRF DRTRICRPPO EISQDGTSYS AOLAGPNAKE WRATGDWAKV TVPLDALRTR TEAWQSYMAV aggageetge ccaccttgtc agaagatcat agcagacgcc gggtgaagaa aggtctgagc ccgctcctgc gatgcaggac cagactccgc tggacaggcc gctgcctgct gcagccagct cagocotood RWLDACLAGS REACGPAGRT GPFFGGAACQ LILRRCELDE KVISVTVKPP SVWRYIRSER TRECNGPSYG aaaaa ccggaaaagc MRGOAAAPGP FPANASRCSW ASCSQGRQQR LGPWSWRGCR ccctcgggaa AGGPENCLTS CSSTCGRGFR GVSEVIOTLV ATDISFPMKG LORNTTVLNS NYCWLSLEGG EEKLKLAHAK aaccggaagc gaagaagcag GVLEEGROCN GSQRRERVCS AVRCPRNATG AEENRDKWEE TLIMIVITYV DROEEGNGDS ggggatcccg gagaatgtcg agcgtggagt ctccagaccg gctgctccgc cccgcacccc accagagcca gaaggtgcct ctgtggaccg ctgcggagga gtggaggca PSRAACOMIC SPWSVCSSTC FFLSSFCWVL ctggactttg acgeceacgt aaacccaaa NP_001693.1

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Angiogenesis Inhibitor 1

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cataggaaat agagcgggag PLVGQDIIDL RKLOHAAEKD VEWERSGATI decdededdd

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ccgccgacac cgcaccgtgc tcagacctgg	aagcogtgga agccctgggg	aaatotatga gcagcagoct ccacgctgga		I.1 MTPACPLLLS DPTKYSLYLR AEAAAGLELC	SSQFTCGVLC PGGPAPPAEA	EEWSPWSVCS WSLCSRSCGR TSCANGTQQR	TGWQRRFRMC IYNKCPPNAS EGMSQVVRSL	VDAENKEKWD AVSSDITFPM PPGPGHSHQR	PAEPLITVEL TFAVLAQPPK CLSILASNIL	LIGIIVENKL SLWSSCVVLP VKCOMGVCRA	RLSIDEDEEP RQLDITWLRP EGYPSFLSVD
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Homo sapiens

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AQGKRRSSID

FTVPTIVVED

LVVSFSSLRA

OVGROADRRA

AIALFQTLAV IYDCLMGFPV

SSVAMGVICT

PTGLVTTIVE

aaatgaaggt ttaattggag gtaactatga ataccacgag agacagacat caagcgggtc ccagaggcag cggacccaat gggagtggas agtgatgtga cagggtaatg acccagtgcc gcctatggag acaataacco caaagatttt acctcgacca ggaacccctg ccaaattttg ggctgatgga gttcatcagt ggccaatcta aaaatgcatc taaaatatac gacagtcaag ctgctgcagt aaatattcca tatctttqaa ttatgcccat aaggaaaat acaccctcaa ctgctggaat acacttgtga tcatctctgt acttttctat aatcagtact EESDDG RYRADLKAVR EKCMALMAND gagggacaa aaagtgacat tggaacactt aaaatagatg gatgtgcgac aacaatcctg tgcagcagat acagtcatct gatggagcag tggaagcagg agattctage tgaggtttat gtctggaaca gcaaacata gaggtatett gtgcaagaag caaagtagaa ttctgtttct

in- AB018301 1 5r 58

19501 G Protein-Coupled Receptor KIAA0758

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BAA34478.1

Receptor KIAA0758 Coupled

G Protein-19501

LQSDSSIVTM

HEISSSPGSL

DLSISIDKAE

MAKALIKSPS ODEMLPTYLK

WKVLQQQWTN

VEPYEDLWGN NTTMPFRISM

SHPETYQORF

SLVMTTTVSH

VILGKPVLNT

MMTHVLSTVN VOMSSTVIKS

STVPTQVNSE SPPLSFSOTN

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TSFSILMSPD

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LFYRLVFILH

FEWMLTLGLM

VCWLNWEDTK

OPREVYTRKN

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Ls21632

G Protein-Receptor Coupled 21632

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Receptor GPR64

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HQDVMKL	HODVMKL ACDDIRVNVV YGLIVIISAI GLDSLLISFS YLLILKTVLG LTREAQAKAF	YGLIVIISAI	GLDSLLISFS	YLLILKTVLG	LTREAGAKAF	
VSHVCAV	VSHVCAV FIFYVPFIGL SMYHRFSKRR DSPLPVILAN IYLLVPPVLN PIVYGVKTKE	SMYHRESKRR	DSPLPVILAN	IYLLVPPVLN	PIVYGVKTKE	
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Leukotriene B4 Receptor BLTR2	Cadherin EGF NM_O LAG Seven- Pass G-Type Receptor 1 (CELSR1/Flam ingo)	

cgactactgc catcacggac ctgtgaggtg catgittgag gattgtggaa gtcggctccg ccgcctgtg cagtttccc cctcaactac gggcgaactg ggtgtctgtg gaagttcctq caccaaggac cctqaacgtg gcgcgtgctg ctgccgcagc cctgagacag cacaggagag cgatggggac ccgggagaat tccccttagc ggtggcaaag tgccatggtg ggaggacctc gtgcgtgtcc ccagagaggg cctagagatc ccagggttcc cagtgtgatt tgtcagtgag ctacacdctc ggáccgggac caacgtcac gccctactd ttcgtctgaa acaagtccaa cgctcatgga ceteccate tctccacgca actacatgaa aaccgctttg cactcagcag acacagtgag accaddtcdc acaccaccac gggatttcta gcagtcccac tcatgtatca tgcaggccac agaatgacaa gtgtcaccat gtcccagga gctccaacat ccaccgtgct gcttcaccgg ctggagagca gtgacgccaa aggagcagca atgtcctaat acgatgagga agttccgcat tctctgccac ggacgacgg gccggctgga acaatgcccc rggggtcggt gggacctgcg tgtcagacag acccdccac gctgtccac ccaacggccg agtatgagag aaaggaacgg tggagatcgt ctcgtggacc accactcagg acctacqaqc gcccgcgacc gactacaage cacactgcgc agctcccatt ctcagtgcca cccgtgccgc gactatgaga cagaaatcag cagttcctgt atcctccagg cgcacccago gtggatcggg gacattaatg aacaaccag aatgcccaga ctgctcaacg gtgctggtgg tatgtcacca gacccgacg ccgctggagg tgcaccctgc ctggagaaca gradecaeca accgacgtca cgcggccagt ctgaccacca ccctgcgaga ctcagctcca tgcccgcccg ccgtgcggcg gaggacttca gtgtgcaaga cctcctggcg ttcatcgccc ttccagggtg ttgctgctgg gaagcacgac cttcaacaac gegetgeege ctactccgac cacdcadccc caacaccgg gctgtacacc cggtgtgatt tgttgaggag tgaaggccct ccagctggac gegggagtat gcacatcctt cactgtccgc cgtggagggg ccagaacgac tggcggcgtc teggacgetg cctgcgcgag cgcgcccttc cgagtgcttc gccaacggg cccdcccag cctcacgttt gaccetgeag gctacctctg cacacggtcg tgtctttcag cattgctacc gattcaggac gatggagctg cggcatcccg caatgcacc ctcgaccagc ggctctggct gaccatcttq cccggcccat gatgagaatg caacaaccgg cacggccttc ctgcgtgtgt gctacacctg geggetteca tcaccatctc gcttcaatga tggagctgtt acgaccctga gcgacctgga ccaacagcat tggccctctt tcttcaacgt cgctgctgcc tctacctgaa acaacatctg tegacetetg ccaggagett cccacaggcc tgtacaccat cccaggacaa atgccaatga atgetecace agcccacgtc ggcatttctt ttgaggtccg gagccacggt tccagatcct teggetgeat agggcaacga tccacagcgt tegacagete tcaacggcct caggccgctg gcagcgtgct tcacaggcgg tcaccctggc catccgacgg tgggcacctc atgggcgtct acaacctttg aaatccaggt tcacctacgt tacaacggcc gcctcggtag attegtgeta gggacatgc gagctggact ctggtgagcc ctgcccgact accggcgtga accttcgtgc cageteagee tctgatggca gacatgctga teceegetge caggagcaga cgcttccact acctaccage ggcggcctca atctttgagg tcaggtccca gtggccgtgt aaggacgaac gacgtetteg gttctgcgat atccaccca gagacggaga cgcgagggcg ctdctcatcd gaggtgacca gccgtgggga gcggtgacag gatgccaaca gacaggcctg aatgcccgca agtggcacca accatcatgg ctcatcctcq ttctacatcg accttctcgg ccttcgacg gatgcccgct

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Cadherin EGF NP_055061.1 CELSR1/Flam Pass G-Type LAG Seven-Receptor 1 ingo) 73584

ROCHRCDNPF KHIVTMTIDY CHINPCENMG SGEKGWLPPE **EAVLMDISRR** GLOGPEVLLE DADSGENARL PVPOFRIDED **OFLWDFYQGS** VLVVOATSAP DPDVSDSLNY CTLRVTITD TDVSSNILNV PCENYMKCVS PCGANGRCRS PPGEYERPYC FIALEIVDEO GERMAVVTVD YLCECPLRFG ATSGGPTSFR INVATLNMNN SKGFDPDCNK EEFPRELESS ERPVLVEFAI MRFYYVVGWG VLSAKVSCOF LRTDLGESTA LSIDEOSSSY SDSEDPSGKE NKVTYPPPLI ARDRDANSVI RTORRLDREN NNPVGSVVAK PVHNRQFVGC NDVRTAYOLI GGTAQLLRRI DDAGOFAVAI HTAHVLINV **QCACKPGVIG** TRPGPGTERE APISRRRHP NTTFGDGPDM YNGRENEKHD GGTCVNRWNM LKNVKEDSEM MOGVRMGGTP ATOHTGTLFG GSALLAPATR AAWEQIQRSE DIFDKFNFTG ARVPRFDTIH VACOCSHTAS HSIHKHLAVA TEVRNIDIGP AVIIINTVTS SFHYLFALFS PGHDSDSDSE AGWPDOSLAE GYPVVHIOAV AVGSSVLTLO AVTASDGTRS KDELELFVEE DVEVENVOND ETEIDICYSD LLIGGEHCVC GHLGLPHGPS GGVPNLPEDF PRKEDSVLME GYLGINCVDA PVCGPCHCAV GSVGNAVRHC SEGAPLPRPL NARITYVIOD FYLEPTSGVI TGVIGCIPAH SDGIHSVTAF PFDDNICLRE VCAELDREEV LILDANDNAP ELDFEVRREY CELLSRNRTH HSRTCDMATG VESLHVYRML LIWSFAGPIG GLLAVNRDAL ATLITESINC KGDAVANHVP GGAARLASSO TYELRINEDA DYKOEQOYVL DGEWHHLLIE ALQLVRALRS NTPMVSTLVY SLVRMLRSNL SIMPRSCKDP OATVLENVPL **DIHNSSGMIT** DINDNAPMFE CPPGFTGDYC VCKNGGTCVN VSVRRGFRGC WEDYSCVCDK LPCPRGWWGN LSANDEDTGE FOGGDDGDGD LINGDLRAMV YVTNKSNSFP PLEALMEVSV VAAVLSTTKD LTTISTORVL ATOERNGLLL QVQYYNKPNI NECDGRRCON VPWYLGLMFR GOPAAVPCPK OKSDITILEI SLDLTGPLLL REHETISLTE GTREGCAARR CPPNSRCHDA LPCDCFPHGS AGIWWPOTKF RNETQVDGAR GGTGGWSARG LILISATWLL VRGSHGEPDA RGEYPPDQES IHPINGLRCR SMSDINIIS SVMLSGLRVT YGPYCENKLD ADFHEDVIHS IVIANMILAV IYMSTEAWTL DECWLSLODT PARGAVHSTP NPAPTPDFPF NDNDPVFTOP DRPVGTSIAT GDMRHFFQLD SPLLALFVEG DARSGRCANG GVSDGRWHSV AQGTQTGSKK VVGGASEDK SLRLPHRPII AALLVAFVLL NEPIEVSSPE GGLITLALPL TIMAQDNGIP SGPNGRLLYT ASVEIQVTIL LPDFOILFNN QLSRDLDNNR **DEQIYLNRTL** PAGRRTTPOT NAAIHYSILS VSVQVLDVND FLGGGSAGPK TSVSITVLDV ILOVSATDRD VDRGSPTPLS VDMAGFIANN EVSHGPSDVE DVDDPCTSSP YYKLLAQDTC IYNGCPKAFE DLRAMNEKLS OGFDLAATOD VRRTYLRPFV PEEKEGPLLR CTVVAILLHY GLDPQGYGNP VSLLRTAFLL HLKGVLGGRK IOKLGVSSGL DGVGAEEKWD ELHREEQGSH NRFALSSORG SSHYTVSVSE DYENOVAYTL SEVIERGLRO GKDIGNYSCA POLFSGESVV GMLPGLTVRS GYVCECGPSH LPERYDPDRR CVEWNHSLAV VTYAAVSLSL NAQIMYQIVE LVDQNDNPPV LENMSQEKFL RGQFFPSEDL LSSTTVLFRP EDFTGEHCEV TTTVAPKVPS LLLDPATGEL TYQLTGGNTR VIIYRTLGQL LEVEERTKPV HCVLNQEVRK /QATDRDQGQ RPPLINSSGV DHGSPPMSSS FEDAPPSTS **IRANDPDEGP** LVSRATVHIL TFVQGNELRL DMLTNSITVR VLREDSSAPE DCDTTMAVRE MRNLSVDGKN COLLINYLOF GMDQNKADIG ALKVRVKDGC INGOCOCKEN AEVITIGCEV LENCTTISEV GHVLQHESWQ EGYFSNVARN VSFPADFFRP ENGEVLPLKI GINOTENPEL IPAIVTGLAV KHHYYGKKGI SLDSIVRDEG ASSHSSDSED RLKVETKVSV HYRLVDTAST DANTHRPVFQ SGIMYIMMEL VAVYNIWALA REGGYTCECF EVITRSEPPO GKNCEQAMPH ACVRSPGSPQ *PESALLPGGV* **JOLT FSAGET**

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ctgcaacttt aagtacagag cacagaagca tacaccgaga MGSPWNGSDG MLIGRYRDMR CTYATLILHWT CDDGISVVPG	IVVVLAFIIC KYRAAAFKIL atggacctgc ccgctcaacg agcctggtct ctgaaggcgg	grettegegg agtgeaggec tgeattect gtetttgggt aacacacegg ceggeeeget	oggacacy oggacacy aacgactaca ggactata aagggtcata taa MDLPPQLSFG	LKAVEALASG CYSWGVCAAI PARFSLSLLL NASNVASFLY atgcacaccg	yycrycugu gaegectgge tegetggtea ctgetgtaec atcageagg tegtaegtga getgteage
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160055 Motilin Receptor (GPR38)	160059 G Protein- coupled Receptor GPR40		160059 G Protein-	coupled Receptor GPR40 160189 G Protein-	Receptor GPR54
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aacgccac agcaacac	gergerggge	ctgccctt	gcaaact	gccarc	accgtggc	rgcccta	acaatgt	gacat	cgago	ttcgtgcg	tgttcag	ნნნანა		gaga	:: ::	gggccc	gaaat	cactcaca	cgcagt	tct	ctgtgaat	ctgaga	gaggcc	tcggaagg	gga	gagaa	ນຂວນ	gctcaggg	taacccta	tcgagg	gcagcag	gggaaa	acagcaca	Ο.	ggctgg	tcacttcc	tgtgt	tggatgaaat	

Receptor GPR44 (CRTH2)

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tttctgccac caaaggccag ggaacaftga ggtgcccagc coctcccatc cottccccct tgcttgttta ttatgttttc gtctattgtc tgtatttgcc				atgaatgaat ccaggtggac	ycytccyayc yccactccry atottcgaga cagtggttat gttatctttg cotttcattg				gootatitia ciggoittat ticacotaci tocacatiti agagoogat tocatatia			MNESEWTEWR ILNMSSGIVN VIFAFHGAPL LHHYTTSYFI ISVLKSVSMA CLACISVDRY GKPGYHGDIF EWCATSWLTS RARFPSHEVD SSRETGHSPD LITTWLAVSNS FCNCVIYSLS	atgagtcagc aaaacaccag accctacagt ttgcagtcca
	NP_004769.1			NM_005684								NP_005675.1	NM_005683
	160210 G Protein- Counled	Receptor GPR44	(CRTH2)	160212 G Protein- Compled	Receptor GPR52							160212 G Protein- Coupled Receptor GPR52	160217 G Protein- Coupled
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	SEGAF	LLDVL	ATEQVHTATM	EALESWKDMN	VETVDNLLRP	KNYNKMHKRE RICKDYIKAV VETVDNILRP EALESWKDWN ATEQVHTATM LLDVLEEGAF	KNYNKMHKRE
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sapiens	PGTYK	FPDPC	QCVVVAGSDA	IMSQRCNNRT	VQCYLPDAFK	ANYGRIDDKI CDADPEQMEN VQCYLPDAFK IMSQRCNNRT QCVVVAGSDA FPDPCPGTYK	ANYGRIDDKI
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Homo sapiens

DRNTIHKNIC VSEVIVVNLV KTSAMRSNTR LOPRGGTSPY SYSLRSGDFP PPVPGGGGEE TSRPLSSPPG LLVEVFESEY INKESVVMAY LVARNPLOGY ENATVKLAGE AGPGGPGGAS LVVNSQVIAA EIYYTSRPPA CRLVESNKTH TECELRGIQT NYFIWSFIGP GLTWAFGLLF SPPGGTHGSL MGNHLLTNPV LCLEGVHLYL TLPLNGNFNN KGPPPPEPPV SESCIAEDGA TEKACWLRVD PPPPAPPGPP SMLGYWSTQG LGGREACGMD NNLRGSSSAA SIVCLAICIS HYFFLAAFSW CLRHSYCCIR SVLYQSDLDE LGAIALLFLL NSTPTLNRGT NCSFWNYSER AAAIDYRSYG SRLDNIKSWA **OKKVHKEYSK** PGSYREPKHP EKMIISELVH EEPLLLPRAQ SPEGPSEALP GPDGDGQMQL SVITWVGIVI TESSEMAGDI HLEDKNHFNA YNNLGLELST LVGIDKTOYE YCFPALVVGI RSSSVLKPDS NESSPEVENS LRDSPSYPDS VFI FVFHCAL RMWNDTVRKO GRNLADAAAF YLAAPGLEGP AEIELLYKAL **IGVVKVVFIL** LMDPVIFTVA LFTTFNAFQG YYTGTQSRIR NTLIAESVGF PGDGGPEPPR EAGGPGGADR YQVRRPSHEG INLFLAELLE SRTKYYYLGG RDSLYASGAN PAVLMAHREI TIMVTIHKMI

gagctgggag ctgctgctgc tgtcgttcct ctctqtccat ttcagggtca agcctcaggg actgagctga ctgggcattg agggcagggg gagtacacca gtcactggtg atcttggtta gatgcccctc ggtcctcgga agcatcatga gatgtggtga tgcttaaagg ggcacccctg cctgaagtct cgggaagagg cccagttta ccagtactcc acageeee catgggctgg gtgtccagaa ctgtcgcctc ccccgggctc tacactcacc cctgtggat tgctggtgac agtggggcc ggatgcgggc gaatgtaaat cacccacgtc gtacaaggag gggcagccc ccaggggcc acaggtcctd caccaaccaa geodeodeod ggagcacccg ccagccagca aggtcgactg cctggaccca cacggatggt togggacccg tgataatgcc ggtgcactat tggagctctg ttgacccggc teccaaegee gtgccctggc ctgtcagggc gtcggaaag aggatgacaa aggatgtgac atgcccagac tgggagacca cctgcgccc tgagggtttg gccctggag cactgcccga tggcacaggc Lgccggagaa agggtgaggc agttcttctc agaccaagag agcagcagga aggggtctgg gaacccgtgg gcaatgccgt tcgacgctga gccgctgcag gtgaccaggg acaccctacq tggtgacagt tcctgcaage ctgggtgggc ctggatcgtg ccccdacdaa ggggtgatcc gtagaggcaa cttctgtgg gacaagggga accaaggagt gtccaggcta ggcgtcccc tcttcggggg cacgatggcc caggccacag gacccggacg cgctccaacc gaggtgctca ccaccactat ctctacacca cctgaaggct ggcaagctca deedeedeed agaagagtcc gagggccatc tattctgtac gqtacccac cagatgccag cccagctac ctttgatage agccgaggag ggttggctat cgctgttttc ccggggacag ctatgagacg catcttcqtc ggttctccat cccggccacc gggacgaggc gascetetgg accaccaget ttccccacad gaccatgac acatagatat ctatgtggtc ctcggatcga aggaggagcc ccacggcatg ccagctgacg gctgctgccg cagtaaccac gtggcaatgc taggctacct taggagccgg agatgcggag tggggtccag gaggccacct ctccacggct aaaggtcacc agttccagcc cggcgcagga ctgacaccaa agaacctgga ccaatgccaa ttgagatcga ggaatccta gtaccacago gtgagaagcg gagtcacage gecetettga acaatgcccc cctcagcgtc ctggccacct atattccct ttgcatccct tggätgccct tgttgctgct

160390 Cadherin EGF NM 001408 LAG Seven-Pass G-Type Receptor 2 (CELSR2)

agctttgggg cttgagcggc tatacagtga tcagatagtc ggggccaca cegeactice tgcaagaatg caccatcaac aatggcacag agcgtgactg gtgcggctca cqtqatqctc tccatcacca cagattgtgg acggatgagg cagttccgca gaccaagtgt gacaccacct atctcagcca ggcgacgatg cggaggctgg gggatgcccc aatccccctg gggctagccg atgtaccaga gagctgacag aatgacaacc cgctcaagca gcctccacgo gtgaccatca tcacccgago ctggccacgc acggccatct tctggagact cgagactcct caggccacgt atcatgagco gactactact tgatttctac gagctcccac cadcatccc ggccagtgtc agctgtggac aaaccgcttc ggactacaaa ggacacggca gatcagcgcc ccagaagtcc cadccccatt cttctccggg cctggtcatc ccttgaccgc ccctgatatc cctgctcaat tctggaggcc cgcgctgcgt ggaggacatg ggccgccacd cagcctgctg gcgggagccc gcccttcatc ctcgcggccc cgattgccca agagtacaca ggactacgaa cagttcctq cgtcctgcag cttccaagga gcgaacgcta tgtgaatgac tgcccagatt totcaccaat cgacgcccc 688886888 ctgccgctgc cccgggtgtc agtggacaag tcactgcctc tgacggtgtc acttcccctt gggaggaagt ttacccaacc tcatggagga atggcattcc tcttctacac gggcatatgc tggaagagaa agcgggacac acctcaaccg actectede gccgttgcac gcaatactcg ccctdccact gcactcggca ctqtcttca cggtggtgct acaatdccc ccttcactag caggcatcgt ctgtgttgga aaggcaccaa agctggacat ctgagtacgt acgtccgcct tcaacaacta ctgcccatga tcagcctggt acaaccggcc ccgcccagtg tccaggcggt cgccagggcc acatctgcct gaggactgcg acctctgcta acacctgcct gttcaagtg aggctgagct cgctgcgcct ccttgctggg gtgggacatg tatgtcttgc gctacagtcc gaactggacc actccagcac gaggtettee gagateett ggaaatgaac aatccaacct accagcgtgg gtatcccttg acccatcgtc gcaggcacca gtcaccaccc gctcgggaca gacgtgaatg gatgtgccac aatggcaggg gatgtgtttg gaggaccggc ggccgagtac gcactggaca cacagcgtga cacagcatca ggcctcttca ttcaacgtac gtgggccagc gagcgcctat ttcgacgaca accgaggtgg gctcgctcag ctggtgggcg atcaccagtg gcctccgatg atcacctact gagtccacgt gtgacagtca gaccccgatd ctgcgcttcg caccccgtcg agaccatggc caacdacaac agctgtgggc cacctaccag cgacgccaac gaatgcccgc cctggtgaac tgtctatgag cgtggcccag cttagactac ggtgagccgg ctttgagcgg ctactgcgag tggtgggctg ggctgttacc ctttattgtt ggatgagttt caacatccct agacggcgta gatgctcacc cgtggtggtc cgtgctgccc ccqqcccatc ccgcagccgc tgtcaacctg tgtggctgct ggaccggccg cacgggggct ggccattact ttctggactt acctatggaa cacagccact gggcaacttt gggtgccatt gctaagccgc accactgcta gagcctgtcg ggacctgcag cgtgtcggtg tgaggtgagt ggaataccd gctggatctc atgaggatge atagtgtcat tgaatgtcac atgttaatga acacaggtga cgatgcaga acctggagat gagacggtga atcgagagaa tctttgagca gtgagctgaa tcaccgatga getteetgte gggcacctg tagaageteg cctggatgt scaaagtgg agtatgtgt cttacacct accagggcag ctgatcgtga cagcccgcac tggcccgggt ttgtggaggg ccctggtaga cagctcctct caccagtgct gettecetgg gacttacad gctggtgtc saccggacca tcctcaacgt cgccctctga cggcacagcg cacgggtga gtgagcactg acatgogotg acgtgatatt acgggcgctg

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160390 Cadherin EGF NP_001399.1 LAG Seven-

Pass G-Type Receptor

(CELSR2)

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SIVILINASTG

TYSFERGNEL TDEMLTHSIT LNVSLSVGOP MRCVSVLRFD GRCRSREGGY EKPYCOVITR VIQEQVQLTF WTVDGCDTG **QEVGCMRNLQ** PLGFGGKSCA RSTITLQLRE YGQQRAEGNL HGESINVEOG SVCTRKPSAP KTSGECHCKE FAEVTINGCE NLENCTSITE ATRLLAHEST

AHDPDISDSL

FPGGAIGRVP LVSDGVHSVT

NNYVTNRSSS NRPLEAIMSV

PVIGNFEILF

ELKLSRALDN

FLSPLLGLFI

PDHVVVFNVQ AQRVL.P FDDN TGDYCETEVD

LNRSLLTAIS

QAVAATLATP

AQCALRVTII RDTDAPGGHI

VRLLDRNDNE

EYVLVIQATS APLVSRATVH

/EGNIPEVFQ IDIFSGELTA LVDLDYEDRP

SSAPFIASSS TCLCRDGYTG SFPAHSFITF SAGESTITVS VALREGSVLG VDSRHIDMAD

CLREPCENY LCYSRPCGPH NEKHDEVALE POGPSEOKVA LPESFPVRMR NOWDAFSCEC

NKPLLGQTGL GPLLLGGVPD NTCHNGGTCV

DGLLLYNGRE

GTCVNLLVGG

RCTPGVCKNG

LALSFATKER

GGSKKSLDLT CPAKKNVCDS SLPISOPWYL PGRANDGDWH AGGVARGFRG DWDSYSCSCD

> FIANNGTVPG GSSLVAWHGL GLQASSIRLE NITVGGIPGP **PCPANSYCSN**

NYSCAAQGTQ

ZWHTVQLKYY

GLRCRCPPGF

VLFRPIHPVG

EHCEVSARSG

RGLRORFHFT

PEVPGGVSDG

PSEDLQERLY

FKCDCPSGDF

PGPGGGPPFL

GPRLHGLHLS

HGYTCECPPN NHYRPPGSPT VNYDSCPRAI

CSIPDFCDSN

PEGVNSLDPS

VCDLNPCEHO VSKGFDPDCN

GVLLOAITRG GPGHAILSFD

HAQLALGASG

CLQGVRVSDT PGYYGDNCTN HPTCGPCNCD

SIMFRTROAD

GROCDRCDNP

GOCPCKPGVI KGSFGTAVRH

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SpeciesName	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens
Peptide	CAPASFERKNERNAEAKRKM	GRIFRAARFRIRKTVKKVE	RTPEDRSDPDACTISK	RHGASPAPQPKKSVNGE	KOTPNRTGKRLTRAQLITD	SPGSTSSVTSINSRVPD	KVRVSDALLEKKKLMA	ANLSSAPSQNCSAKD	IKLADSALERKRISAA	GEASNRSLNATETSEA	RIYRAARNRILNPPSL	KAQEEMSDCLVNTSQIS	RHLSNRSTDSQNSFASC	CTTEASMAIRPKTITEKM	DNDLDHPGERQQISST	CVSDFSTSDPTTEFEK	RIYHAAKSLYQKRGSSR	ESGEKSTKSVSTSYVL	DKCKISEEMSNFLAWLG	IAKEEVNGQVLLESGE	STVRSLRSEFKHEKSWR	DAFNWTVDSENRINLSC	FGLQDDSKVFKEGSC	PGSYTGRRTMQSISNEQKAC	CSMVALGKQHSEEASKDNSD	NTIPALAYKSSQLQMGQ	KGIETDVDNPNNITC	CSSPEKVAMLDGSRKDKA .	RRTSTIGKKSVQTISNE	CNYRATKSVKTLRKRSSK	SGLQTESIPEEMKQIVEEQG	CKRNTAEEENSANPNODONA	GHTEEPPGLSLDFLKC	CNYKVEKKPPVROIPRV	IGLRDEEKVFVNNTTC
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Source ID	P08908	P08908	P08908	P08908	P28222	P28222	P28222	P28222	P28221	P28221	P28221	P28221	P28566	P28566	P28566	P28566	P28566	P30939	P30939	P30939	P30939	CAA01675.1	CAA01675.1	CAA01675.1	CAA01675.1	CAA01675.1	P41595	P41595	P41595	P41595	P41595	P28335	P28335	P28335	P28335
Gene	5-HT1A Receptor	5-HT1A Receptor	5-HT1A Receptor	5-HT1A Receptor	5-HT1B Receptor	5-HT1B Receptor	5-HT1B Receptor	5-HT1B Receptor	5-HT1D Receptor	5-HT1D Receptor	5-HT1D Receptor	5-HT1D Receptor	5-HT1E Receptor	5-HT1E Receptor	5-HT1E Receptor	5-HT1E Receptor	5-HT1E Receptor	5-HT1F Receptor	5-HT1F Receptor	5-HT1F Receptor	5-HT1F Receptor	5-HT2A Receptor	5-HT2A Receptor	5-HT2A Receptor	5-HT2A Receptor	5-HT2A Receptor	5-HT2B Receptor	5-HT2B Receptor	5-HT2B Receptor	5-HT2B Receptor	5-HT2B Receptor	5-HT2C Receptor	5-HT2C Receptor	5-HT2C Receptor	5-HT2C Receptor
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	RHTNEPVIEKASDNEP RNAVHSFLYHLIGLLVWQCD	CDISVSPVAAIVTDIFNTSD	DGGRFKFPDGVQNWPALS	NNIGIIDLEKRKFNQ	ESRPOSADQHSTHRMR	CDDERYRRPSILGQTVP	RDAVECGGQWESQCHPPATS	VTAKEHAHQIQMLQRAGASSESRP	KSFRRAFUILCCDDE	VTAKEHAHQIQMLQRAGA	KEHAHQIQMLQRAGA	VIAKEHAHQIQMLQR	RTPRPGVESADSRRLATK	CPRERGASLASPSLRTS	PLFMRDFKRALGRFLPC	RAAAAVNFFNIDPAEPE	EVTASPAPTWDAPPDNASGC	KAARKSAAKHKFPGFPRVE	CANLSRLIKHERKNISIFKR	KLAERPERPEFVLRAC	CHKPSILTYIAIFLT	NGSMGEPVIKCEFEKVISME	NKKVSASSGDPQKYYGKELK	NDHFRCQPAPPIDEDLPEER	CQPKPPIDEDLPEEKAED	QPKPPIDEDLPEEKAED	MPPSISAFQAAYIGIEVU	GENIGLPDVELLSHELKGVC	MPIMGSSVYITVELAIA	RSHVLRQQEPFKAAGT .	RIREFROTFRKIIRSH	KDSATNNCTEPWDGTTNES	CROLORTELMDHSRTTLQRE	RNRDFRYTFHKIISRYLLC	CQADVKSGNGQAGVQP
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RSNRRGPRAKGGPGQGE	ASAREVNGHSKSTGEK	RGVGAIGGQWWRRRAH	RAPVGPDGASPTTENG	RTGTARPRPPTWSRTR	ASRSPGPGGRLSRASS	RSVEFFLSRRRARSSVC	PMASGRQQRRRQARVTC	NYHILASLRTREEVSR	RVRGPKDSKTTALLT	VGRLFRTKVWELYKQC	FRTMKEYSDEGHNVTAC	CIMOIMQVLRINNEMOKFKE	CODERIIDVITQIASFM	CRSEPIQMENSMGTLRTS	RVFREAGKGVKKIDSC	CERRFLGGPARPPSPS	ANGRAGKRRPSRLVALRE	CARRAARRHATHGDRPRAS	CLARPGPPSPGAASD	CNGGAAADSDSSLDEP	KRQLQKIDKSEGRFHV	GEOSGYHVEOEKENKLLC	APNRSHAPDHDVTQQR	VPLVIMVFVYSRVFQE	RGELGRFPPEESPPAP	SRSLAPAPVGTCAPPE	GVPACGRRPARLLPLRE	PSGVPAARSSPAQPRLC	EEEFYLFKNISSVGPWDGPQ	CGPDWYTVGTKYRSESYT	NNRNHGLDLRLVTIPS	IMKMVCGKAMTDESDT	SITNDTESSSVVSNDNTNK		KAVVKPLERQPSNAILKTC	
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P18089	P18089	P18089	P18825	P18825	P18825	P18825	P46663	P46663	P46663	P46663	AAB02793.1	AAB02793.1	AAB02793.1	AAB02793.1	AAA51667.1	AAA51667.1	AAA51667.1	AAA51667.1	AAA51667.1	AAA51667.1	NP_000015.1	NP_000015.1	NP_000015.1	NP_000015.1	P13945	P13945	P13945	P13945	NP_001699.1	NP_001699.1	NP_001699.1	NP_001699.1	AAA35604.1		AAA35604.1	
Alpha 2b-adrenoceptor	Alpha 2b-adrenoceptor	Alpha 2b-adrenoceptor	Alpha 2c-adrenoceptor	Alpha 2c-adrenoceptor	Alpha 2c-adrenoceptor	Alpha 2c-adrenoceptor	Bradykinin B1 Receptor	Bradykinin B1 Receptor	Bradykinin B1 Receptor	Bradykinin B1 Receptor	Bradykinin B2 Receptor	Bradykinin B2 Receptor	Bradykinin B2 Receptor	Bradykinin B2 Receptor	Beta-1 adrenoceptor	Beta-2 adrenoceptor	Beta-2 adrenoceptor	Beta-2 adrenoceptor	Beta-2 adrenoceptor	Beta-3 adrenoceptor	Beta-3 adrenoceptor	Beta-3 adrenoceptor	Beta-3 adrenoceptor	Opsin, blue-sensitive	Opsin, blue-sensitive	Opsin, blue-sensitive	Opsin, blue-sensitive	Bombesin Receptor	Subtype-3	Bombesin Receptor	Subtype-3					
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Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Mus musculus	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens
RDPNKNMTFESCTSYPVSKK	RTLYKSTLNIPTEEQSHARK	KSFQKHFKAQLFCCKAERPE	NKGWSGDNSPGIEALC	GROPHSPNOTLISITNDTE	RPEPPVADTSLTTLAV	SEISVTSFTGCSVKQAEDR	ELDRIDNYNDTSLVENHLC	SQGHHNNSLPRCTFSQE	CYVGVVHRLRQAQRRP	CQLFPSWRRSSLSESENA	TEDYDTTTEFDYGDATPC	ASMPGLYFSKTQWEFTHHTC	CSLHFPHESLREWKLFQA	TILISVFQDFLFTHEC	CSALYPEDTVYSWRHF	PEFIFYETEELFETLC	SSYQSILFGNDCERSK	GRYIPFLPSEKLERTS	DDVGLLCEKADTRALMAQFV	MNATEVTDTTQDETVYNSYY	DESIYSNYYLYESIPKPC	DIPSSSYTQSTMDHDLHD	LETLVELEVLQDCTFE .	RNHTYCKTKYSLNSTTWK	CODEVIDDYIGDNITVD	PELLYSDLQRSSSEQAMRC	QLRQWSSCRHIRRSSMSVE	GVKFRNDLFKLFKDLGC	PDIFSSPCDAELIQING
22	23	24	2286	2287	2288	2289	1382	1383	1384	1385	305	1242	1243	1244	1386	1387	1388	1389	1751	306	348	351	353	491	748	846	847	848	359
AAA35604.1	AAA35604.1	AAA35604.1	NP_001718.1	NP_001718.1	NP_001718.1	NP_001718.1	5 P32302		5 P32302	5 P32302	P32246	P32246	P32246												P32248	P32248	P32248		P51685
Bombesin Receptor	Bombesin Receptor Subtype-3	Bombesin Receptor Subtype-3	Bombesin Receptor	Bombesin Receptor	Subrype-3 Bombesin Receptor	Bombesin Receptor	CXC Chemokine Receptor 5	C-C Chemokine Receptor 1	C-C Chemokine Receptor 3	C-C Chemokine Receptor 4	C-C Chemokine Receptor 7	C-C Chemokine Receptor 7	C-C Chemokine Receptor 7	C-C Chemokine Receptor 7	C-C Chemokine Receptor 8														
769	692	692	769	692	695	692	729	729	729	729	735	735	735	735	737	737	737	737	737	738	738	738	738	738	74	74	741	74	742
827	828	829	830	831	832	833	834	835	836	837	838	839	840	841	842	843	844	845	846	847	848	846	820	821	825	853	854	855	920

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Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens		Homo sapiens	Homo saplens		Homo saplens		Homo sapiens	Homo sapiens		Homo sapiens		Homo sapiens	Homo sapiens		Homo saplens		Homo sapiens		Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens
KILHQLKRCQNHNKTKAIR SOIENVI CROMADDESC	FVGEKFKKHLSEIFQKSC	ENFSSYDYGENESDSC	CYAHILAVILVSRGQRRURA	MVLEVSDHQVLNDAEVAALL	CPNGRGLQRQPSSSRRD	TEEMGSGDYDSMKEPC	KKLRSMTDKYRLHLSVAD	CIIISKLSHSKGHQKRKALK	KILSKGKRGGHSSVSTE	ENRSLENIVOPPGEMNDRLD		KIPSGFPIEDHETSPLDNSD	RKKARQSIQGILEAAFSEE		PGTFQRPSADSLPRGSARLT		DLNTPVDKTSNTLRVPD	CGVDYSHDKRRERAVAIVRL		CYTFILLRTWSRRATRSTK		GGRLRKSLPSLLRNVLTE	AELEESPEDSIQLGVTR		EFVLIPWRPEGKIAEEV		RRNWNQYKIQFGNSFSNSE		RSASYTVSTISDGPGYSHDC	NDIQYEDIKGDMASKLG	KENEENIQCGENFMDIE	EDGKVQVTRPDQARMDIR
360	493	1371	1372	1373	1374	1376	1377	1380	1381	25		7 0	27		28		811	812		813		814	841		843		844		845	53	30	31
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P51685 P51685	P51685	P49682	P49682	P49682	P49682	P30991	P30991	P30991	P30991	AAC50657.1	. (AAC5000/.	AAC50657.1		AAC50657.1		P21730	P21730		P21730		P21730	Q16602	• •	Q16602		Q16602		Q16602	AAB18200.1	AAB18200.1	AAB18200.1
C-C Chemokine Receptor 8 C-C Chemokine Receptor 8	C-C Chemokine Receptor 8	CXC Chemokine Receptor 3	CXC Chemokine Receptor 4	Complement Component		Complement Component 3a Receptor 1	Complement Component		Complement Component		Complement Component	Complement Component	5a Receptor 1	Complement Component	٠.	Complement Component	Calcitonin Receptor-like		Calcitonin Receptor-like		Calcifonin Receptor-like	Receptor	Calcitonin Receptor-like Receptor	Cannabinoid Receptor 1	Cannabinoid Receptor 1	Cannabinoid Receptor 1						
	742 (755		8	755 C	ຕ	755 (-,	758 (758		758	-	758	767		767))		/9/	832		832 (
857 858	859	980	861	862	863	864	865	998	867	868	070	6	870		871	į	872	873	. 1	874	į	8/2	876		877	0	۵/۵		6/8	880	881	882

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	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens		Homo saplens		Homo saplens		Homo sapiens	Homo sapiens	Homo saplens	Homo saplens					
	DPEGPYSYCNTTLDQIGTCW	ALLEGYCHTIMTLTNLSG	SSHHEPRGSISKEC	KAKPTSPSDGNATSLAETID	CSQPESSFKMSFKRE	EDLKKEEAAGIARPLEK	PWEEDFWEPDVNAENC	CAPDISLRASIKKETK	PNAVTPGNREVDNDEE	QTSPDGDPVAESVWELDC	KRSSRAFRAHLRAPLKGNC	CTVIMKSNGSFPVNRRRV	KPEKNGHAKDHPKIAK	GKTRTSLKTMSRRKLSQQKE	KGRRRKRILTRØNSQC	CNSVRPGFPQQTLSPDP	CQDTALGGPGFQERGGE	KREEKTRNSLSPTIAP	STSLKLGPLQPRGVPLRE	VAVAVPLRYNRQGGSR	EVARRAKLHGRAPRRP	PPSPTPPAPRLPQDPC	PPQTPPQTRRRRAKITGRE	DAYPSAFPSAGANASGP		LVDIDRRDPLVVAALHLC		KRCFRQLCRKPCGRPD		SKRIKEALAKERVIAC	TENSSQLDFEDVWNSS	NDSFPDGDYDANLEAAAPC	CHASLGHRLGAGQVPG
	505	507	41	42	43	4	1407	1408	1409	1410	1403	1404	1405	1406	1398	1399	1400	1401	1402	1394	1395	1396	1397	222		224		225	ò	077	1411	1412	1413
	24		CAA41734.1	CAA41734.1	CAA41734.1	CAA41734.1	8	8	8	. 8	9	9	. 9	9	2	7	20	2	22	7	7	7	7	AAA18789.1		AAA18789.1		AAA18789.1	1 0000	44410/09.1	AAC50055.1	AAC50055.1	AAC50055.1
	Q13324	LR43	CAA	CAA	CAA	CAA	P21918		•	P21918	P14416		P14416	P14416	P35462	P35462	P35462	P35462	P35462	P21917	P21917	P21917	P21917	_							AACE	AACE	AAC.
Idetor Receptor 2	Corticotropin releasing factor Receptor 2	Corticotropin releasing factor Receptor 2	Dopamine Receptor D1	Dopamine Receptor D1	Dopamine Receptor D1	Dopamine Receptor D1	Dopamine Receptor D5	Dopamine Receptor D5	Dopamine Receptor D5	Dopamine Receptor D5	Dopamine Receptor D2	Dopamine Receptor D2	Dopamine Receptor D2	Dopamine Receptor D2	Dopamine Receptor D3	Dopamine Receptor D4	Dopamine Receptor D4	Dopamine Receptor D4	Dopamine Receptor D4	Opiold Receptor, delta	(OPRD1)	Opioid Receptor, delta	(CrkD)	Opioid Receptor, delta	. ₹	OPRD1)	Duffy Antigen	Duffy Antigen	Duffy Antigen				
	1103	1103	1240	1240	1240	1240	1241	1241	1241	1241	1242	1242	1242	1242	1243	1243	1243	1243	1243	1244	1244	1244	1244	1267		1267)	1967) (1)	1424	1424	1424
	915	916	617	918	616	920	921	922	923	924	925	956	927	928	5 26	086	931	932	933	934	935	936	937	338		666	6	940	. [Ī	942	943	44

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Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens		Homo sapiens	•	Homo sapiens	Homo sapiens		Homo sapiens	aclary cmcH	STEPIC OF CHILD	Homo sapiens	Homo saplens	•	Homo saplens		Homo sapiens	Homo sapiens		Homo sapiens	Homo sapiens
FGAKGLKKALGMGPGP	KLFRTAKONPLTEKSGVNKK	KSAPEENSREMTETQM	CKGYKRKVMRMLKRQ	GEERGFPPDRATPLLQTAE	RSLAPAEVPKGDRTAGSP	PRTISPPPCQGPIEIKE	EEKQSLEEKQSCLKFKAND	RYSTNLSNHVDDFTTFRGTE	NRRNGSLRIALSEHLK	EYRGEQHKTCMLNATSK	KNHDONNHNIDRSSHKD	RPGIEKFREEAEERDIC		CHLQEGAKGPLPVDTFLR	. 60	GHEESGDIATSINSSI AFIAPLC	KGIIEGEPTCCFECVECPDG		CSTAAHAFKVAARATLRRSN	BISINGHENDHONDI	י מנו לאולי וונויסנו ומנויסנר	RPEVEDPEELSPALVVSSSQ	ASWGGTPEERLKVAITMLTA		SEDSAPTNDTAANSAS		SYESAGYIVLIRILPLVVL	PVFLFLTTVTIPNGD		EERLKVAITMLTARGIIRFV	ERALSEDSAPTNDTAANSAS
1415 45	§ 4	47	48	22	. 99	99	. 29	49	9	51	53	1425		1426		142/	1428		1429	1/30	<u> </u>	1431	1878		1879	•	1880	1881		2612	2613
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 AAC50055.1	AAA35924.1	AAA35924.1	AAA35924.1	BAA14398.1	BAA14398.1	BAA14398.1	BAA14398.1	AAB25530.1	AAB25530.1	AAB25530.1	AAB25530.1	P41180		P41180		.F41180	P41180		P41180	041180	2	P41180	NP_001453.1		NP_001453.1		NP_001453.1	NP_001453.1		NP_001453.1	NP_001453.1
Duffy Antigen	EBV-Induced Gene 2	EBV-Induced Gene 2	EBV-Induced Gene 2	Endothelin B Receptor	Endothelin B Receptor	Endothelin B Receptor	Endothelin B Receptor	Endothelin A Receptor	Endothelin A Receptor	Endothelin A Receptor	Endothelin A Receptor	Calcium-Sensing Receptor	(CASR)	Calcium-Sensing Receptor	(CASR)	Calcium-sensing receptor (CASP)	Calcium-Sensing Receptor	(CASR)	Calcium-Sensing Receptor	Colci im-Sensing Becentor	(CASR)	Calclum-Sensing Receptor	(CASK) Formyl Peptide Receptor-	Like Receptor	Formyl Peptide Receptor-	Like Receptor	Formyl Peptide Receptor- Like Recentor	Formyl Peptide Receptor-	Like Receptor	Formyl Peptide Receptor-	Like receptor Formyl Peptide Receptor-
1424	1451	1451	1451	1486	1486	1486	1486	1488	1488	1488	1488	1598		1598		0,60	1598		1598	1508	2	1598	1676		1676	į	9/91	1676		1676	1676
945 946	8	948	949	950	951	952	953	85	955	926	957	958		959	č	3	196		962	649	3	8	965	,	99		<u>}</u>	896		696	970

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	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	. Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens
	GESKVTEIPSDLPRNAIELR	DVLEVIEADVFSNLPK	RNGHCSSAPRVTSGSTY	RGQRSSLAEDNESSYSRGFD	CHHRICHCSNRVFLCQE	LRVIQKGAFSGFGDLEK	LYVIMSLLVLNVLAFVVIC	CNKSILRQEVDYMTQARGQR	SDNNNLEELPNDVFHGA	KLVALMEASLTYPSHC	SFESVILWLNKNGIQEIHNC	IHSLQKVLLDIQDNINIHT	KANNLLYIPEAFQNLP	CYEMQAQIYRTETSSTVH	INTPSSRKKMVRRVVC	ARAISASSDQEKHSSRK	KYSAKTGLTKLIDASRVSET	PDTYYLKTVTSASNNETYC	GNSLVITVLARSKPGKPR	PRASNQTFCWEQWPDPRHKK
	58	29	09	61	2231	2232	2233	2234	2236	2238	2241	2248	2250	2251	1437	1439	1440	1893	192	193
	AAA52477.1	AAA52477.1	AAA52477.1	AAA52477.1	NP_000136.1	NP_000136.1	NP_000136.1	NP_000136.1	NP_000136.1	NP_000136.1	NP_000136.1	NP_000136.1	NP_000136.1	NP_000136.1	AAA62370.1	AAA62370.1	AAA62370.1	AAA62370.1	AAA50767.1	AAASU/6/.1
11kg Docostor	nulating Hormone	Receptor Follicle Stimulating Hormone AAA52477.1	Receptor Follicle Stimulating Hormone Recentor	Follicle Stimulating Hormone	Follicle Stimulating Hormone Recentor	mulating Hormone	Follicle Stimulating Hormone	Follicle Stimulating Hormone NP_000136	mulating Hormone	mulating Hormone	mulating Hormone	mulating Hormone	receptor Follicle Stimulating Hormone Receptor	mulating Hormone	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	Galanin Receptor GalR1	Galanin Keceptor Galiki
	1681	1681	1681	1891	1681	1681	1891	1891	1681	1891	1681	1681	1681	1681	1726	1726	1726	1726	1762	70/1
	. 126	972	973	974	975	926	777	876	979	086	186	982	983	984	985	986	786	988	686	5

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Homo saplens	Homo saplens	Homo sapiens		Homo saplens		Homo sapiens		Homo sapiens	-	Homo sapiens	Homos cardens		Homo sapiens		Homo sapiens		Homo sapiens	Homo sapiens	Homo soniens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens		Homo saplens		Homo sapiens		Homo saplens		Homo saplens	Homo sapiens	Homo saplens
KKLKNMSKKSEASKKKTAQ GNSLVITVLARSKP	RKDSHLSDTKENKSRID	QTAGELYQRWERYRREC		CENPEKNEAFLDQRULER		CRLRRSLGEEQRQLPERAFR		PTSRGLSSGTLPGPGNEA		CNISSHSADLFVNDDWSHPG	SPI HPEHEESTNOTEISO		YNLPVEGNIHVKKQIES		CQPGUIRSHSTGRSTT		CEPPRIRGAGTRELELAIR	RVRNQGGLPGAVHQNGRC	LRFDGDSDSDSQSRVR	CRPETGAVGKDSDGCY	DGLLRTRYSQKIGDDL	CGPDGQWVRGPRGQPWRDAS	CQMDGEEIEVQKEVAKMYSS	TSNHRASSSPGHGPPSKE	KLOKWTOKKEKGKKLSRMK		DRSLAITRPLALKSNSKVGG	•	RMIHLADSSGQTKVFSQC		DPHELQLNQSKNNIPRARLK		GRLAGRHPQDSYEDSTQSS	CKPFGNVRFDAKLAIVG	KTSCGPDVFSGSSYPGVQS
194 195	961	1250		1251		1253		1276	Ç G	679	830	· }	831		832		1281	1282	1283	1284	837	838	839	840	206		207		208		506		1746	1747	1748
AAA50767.1 AAA50767.1	AAA50767.1	P48546		P48546		P48546		P48546	030550	Longon	P30550		P30550		P30550		Q16144	Q16144	Q16144	Q16144	P47871	P47871	P47871	P47871	AAA35917.1		AAA35917.1	:	AAA35917.1	•	AAA35917.1		NP_000504.1	NP_000504.1	NP_000504.1
Galanin Receptor GaIR1 Galanin Receptor GaIR1	Galanin Receptor GaIR1	Gastric Inhibitory	Polypeptide Receptor	Gastric Inhibitory	Polypeptide Receptor	Gastric Inhibitory	Polypeptide Receptor	Gastric Inhibitory	Polypeptide Receptor		receptor Gastrin-Releasing Peptide	Receptor	Gastrin-Releasing Peptide	Receptor	Gastrin-Releasing Peptide	Receptor	Cholecystokinin B Receptor	Cholecystokinin B Receptor	Cholecystokinin B Receptor	Cholecystokinin B Receptor	Glucagon Receptor	Glucagon Receptor	Glucagon Receptor	Glucagon Receptor	Gonadotropin-Releasing	Hormone Receptor	Gonadotropin-Releasing	Hormone Receptor	Gonadotropin-Releasing	Hormone Receptor	Gonadotropin-Releasing	Hormone Receptor	Opsin, green-sensitive		Opsin, green-sensitive
1762	1762	1808		1808		1808		1808	1813	2	1813		1813		1813		1814	1814	1814	1814	1834	1834	1834	1834	1925		1925		1925		1925	. 4	1945	1945	1945
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CILQLFGKKVDDGSELSS STRGPFEGPNYHIAPR INGLVLAATMKFKKLR	ELSSASKTEVSSVSSVSP ADLDWDASPGNDSLGD		GVEHENGTDPWDINEC	KLWRRRRGDAVVGASL		SQRKLSTLKDESSRAW	OFDESA O I O A SECURIORITI		CPDFFSHFSSESGAVKRD		VIKLEPAGGSLHIGSQ		KIEISKAVANGADPELL	GWNHFMQQTSVRREDKC	COMPELINISLPSFSEIKLR	AGGGSVLKSPSQTPKE	KSPVVFSQEDDREVDKLYC	TAPGKGKLRSGSNTGLD	KRLRSHSRQYVSGLHMNRE	NSRNETSKGNHTTSKC	CITYYRIFKVARDQAKR	RDQAKRINHISSWKAA	TAFVYRGURGDDAINE	HKTSLRSNASQLSRTQSRE	DSNGSAGSEDAQLEPA .	KVREDVDVIECSLQFPDDD	RNIVQDPAYLRDIDGMNK	CFPLKMRMERQSTSRVRN	
1750 1767 1768	1769 581		582	583		584	833		834	į	835	700	000	1167	1168	1169	1170	1171	1172	1173	1174	1175	1176	1177	227	228	229	230	
NP_000504.1 NP_000504.1 NP_000504.1	NP_000504.1 Q92847		G92847	Q92847		Q92847	502643		Q02643	* XOO	GU2043	000843	QQ7040	P35367	P35367	P35367	P35367	P35367	P35367	P25021	P25021	P25021	P25021	P25021	AAA63906.1	AAA63906.1	AAA63906.1	AAA63906.1	
Opsin, green-sensitive Opsin, green-sensitive Opsin, green-sensitive		ceptor			ceptor		Secretagogue receptor		Growth Hormone-Releasing		eledsing	Hormone Receptor		oto	ō	ក		ō	Histamine H1 Receptor		Histamine H2 Receptor	ö			Opiold Receptor, kappa 1 (OPRK1)	Opiold Receptor, kappa 1	Opiold Receptor, kappa 1	(OPIKKI) Opiold Receptor, kappa 1	
1945 1945 1945	1945 1951		<u>.</u>	1951		1951	1954		1954	7301	<u> </u>	105/1	<u> </u>	2120	2120	2120	2120	2120	2120	2121	2121	2121	2121	2121	2783	2783	2783	2783	
1017 1018 9101	1020	9	770	1023		1024	1025		1026	700	102/	ACO.	3	1029	1030	1031	1032	1033	1034	1035	1036	1037	1038	1039	94	<u>8</u>	1042	1043	٠

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	Homo saplens			Homo sapiens		Homo saplens			Homo saplens			Homo sapiens			Homo sapiens	-	Homo sapiens		Homo saplens	•	Homo sapiens		Homo sapiens	Homo saplens		Homo sapiens	Homo saplens		Homo sapiens	Homo sapiens		Homo saplens	Homo saplens
	CNTGIRKFPDVTKVFSSESN			NVIDINGATIGORILL		CESTVRKVSNKTLYSS			FAVRNPELMATNKDTK			CKRRAELYRRKDFSAYTSN		4	ERHITVFRMQLHTRMSNRR		RGRTMRMSRHSSGPRRNRD		KHLATEWNTVSKLVM		ENPTGPTESSDRSASSLN		ESQISLSCSLCLHSGDQEAQ	QQQKATRVYAVVQISAPM		DKPEVGRNKKAAGIDPME	EQPHSTQHVENLLPREHRVD		RLHVKRIAALPPADGVAPQ	DPLIYAFRSLELRNTFRE	•	QAPFFSNQSSSAFCEQVFI	IVHSDYLIFEDQFIQHMDNI
	1432	•	1,400	554		1434		•	1435		.•	1436	-		210		211	•	212		213		184	185		186	187		451	452		562	563
	Q14751		014761	2/4	·	Q14751			Q14751			Q14751			AAC51139.1		AAC51139.1		AAC51139.1		AAC51139.1	•	AAB21255.1	AAB21255.1		AAB21255.1	AAB21255.1		P41968	P41968		P41968	P41968
(JAKI)	Luteinizing	Hormone/Chorlogonadotro	pin Receptor	Hormone/Chorlogonadotro	pin Receptor	uteinizing	Hormone/Choriogonadotro	pin Receptor	uteinizing	Hormone/Chorlogonadotro	oin keceptor	Luteinizing	Hormone/Chorlogonadotro	pin Receptor	Lysophosphatidic Acid	Receptor Edg2	G Protein-Coupled Receptor MRG	G Protein-Coupled	Receptor MRG	G Protein-Coupled Recentor MRG	G Protein-Coupled	Receptor MRG	Melanocortin 3 Receptor (MC3R)	Melanocortin 3 Receptor	(MC3R)	Melanocortin 3 Receptor (MC3R)	Melanocortin 3 Receptor						
	2964		7064		· u	2964			2964 L	- ! - 1	_	2964 L	·	-	2976 L	<u>.</u>	2976 L	-	2976	_	2976 L		3038	3038	<u> </u>	3038	3038		3057	3057		3057	3057
	1044		1045	} ·		1046		•	1047		:	1048			1049	•	1050		1051		1052		1053	1054		1055	1056		1057	1058		6901	0901

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 Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens							
HSNASESLGKGYSDGGC	KRIAVLPGTGAIRQGA	NSTDTDAQSFTVNIDN	NSTHRGMHTSLHLWNRSSYR	ATEGNLSGPNVKNKSSPC	NKHLVIADAFVRHIDN	MNSSFHLHFLDLNLNAT	RYHHIMTARRSGAIIAG	QGSQRRLGSLNSTPT	EAGALVARAAVLQQLD	ALRYHSIVTLPRARQA	CQHAQGIARLHKRQRP	HSLKYDKLYSSKNSLC	CTARVFFVDSSNDVADR	DSSNDVADRVKWKPSPLMIN	AVRPGWSGAGSARPSR	LVAIFYDGWALGEEHC	LVLQARRKAKPESRLC	CIQDASKGSHAEGLQSPA •	CENTAPORE GLEVISY	APARAHAPINA APANAHAN	DRASGHPKPHSRSSSAY	HPKPAAADNPELSASHC
1032	1033	1035	1469	1022	1024	1025	1026	1036	1038	1039	1040	214	215 216	217	930	931	932	933	754 751	752	753	754
		. •			<i>)</i>		•			 						. :				•		, .
AAB33341.1	AAB33341.1	AAB33341.1	AAB33341.1	P33032	P33032	P33032	P33032	AAD41352.1	AAD41352.1	AAD41352.1	AAD41352.1	•	AAB17/20.1					5 P49286 5 P46584		_	_	or Q13585
(MC3R) Melanocortin 4 Receptor (MC4R)	Melanocortin 4 Receptor	Melanocortin 4 Receptor	Melanocortin 4 Receptor (MC4R)	Melanocortin 5 Receptor	Melanocortin 5 Receptor	Melanocortin 5 Receptor (MC5R)	Melanocortin 5 Receptor	Melanocortin 1 Receptor	Melanocortin i Receptor	Melanocortin 1 Receptor	Melanocortin 1 Receptor (MC1R)	Ф	Melatonin Receptor type (a	9	Melatonin Receptor type 1b	Melatonin Receptor type 1b	Meigtonin Receptor type 1b	Melatonin Receptor type 1b	Melatonia-Related Recentar	Melatonin-Related Receptor	Melatonin-Related Receptor	Melatonin-Related Receptor
3058	3058	3058	3058	3059	3059	3059	3059	3061	3061	3061	3061	3079	3079	3079	3080	3080	0808	2080	308	3081	3081	3081
190	1062	1063	1064	1065	9901	1067	1068	1069	1070	ָ ועסנ	1072	1073	1075	1076	1077	1078	6/0	8 5	280	83	1084	1085

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Homo sapiens	Homo saplens		Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	
DDSDLPESASSPAAGPT	DDYRIGININSGVVRSVC CRSNTFI NIFRRKKAG	DISTUTI VANVEEEED A	Districtive	ERFKLLGEYVYEHERE	DFVRASLSRGADGSRHIC	CVATSEKVGRAMSRAAFEG	CAAHSLRAVPFEQESK	CDAMRPVNGRRLYKDF	DAPFRPADTHNEVRFDR	GKETAPERREVVTLRC	GGLFPINEKGTGTEEC	EFVRASLTKVDEAEYMC	RSNIRKSYDSVIRELL	CDKHLAIDSSNYEQES	GTRRYTLAEKRETVILKC	PSSLGKPKGHPHMNSIRID •	CGSGGPPIITKPERVVG	CKLSRHALKKGSHVKK	CPRMDPVDGTQLLKYI	=
755	880) <u>(</u>	- 00	882	891	892	893	894	895	968	897	868	668	006	902	606	910	911	913	
Q13585	Q13255	013055	Ø13200	Q13255	Ó14416	<u>©</u> 14416	Q14416	Q14416	Q14416	Q14416	CAA54796.1	CAA54796.1	CAA54796.1	CAA54796.1	CAA54796.1	Q14833	Q14833	Q14833	Q14833	
Metabolio-Related Receptor Q13585	Receptor 1 Metabotropic Glutamate	Receptor 1 Metabotropic Glutamate	Receptor 1	Metabotropic Glutamate Receptor 1	Metabotropic Glutamate Receptor 2	Metabotropic Glutamate	Metabotropic Glutamate	Metabotropic Glutamate Receptor 3	Metabotropic Glutamate Receptor 3	Metabotropic Glutamate Receptor 3	Metabotropic Glutamate	Metabotropic Glutamate Recentor 4	Metabotropic Glutamate	Receptor 4 Metabotropic Glutamate	Receptor 4 Metabotropic Glutamate Receptor 4	1 5 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2				
3081	3093	3093	3	3093	3094	3094	3094	3094	3094	3094	3095	3095	3095	3095	3095	3096	3096	3096	3096	
1086	1088	1089	3	0601	1091	1092	1093	1094	1095	1096	1097	1098	1099	0011	1101	1102	1103	1104	1105	

									,										
Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens
RIERMHWPGSGQQLPRSIC	KDYFDYINVGSWDNGEL	KMDDDEVWSKKSNIIRSVC	GETLRYKDRRLAQHKSEIEC	NPNQTAVIKPFPKSTE	KALYDVAEAEEHFPAPA	RSPSPISTLSHRAGSASRTD	RESPAAGPEAAAKPD	QAURGRGDGDEVGVRC	KLTSSGTQSDDSTRKC	DVEALQWSGDPHEVPSSLC	RFQVDEFTCEACPGDM	GARPHSVIDYEEQRT	CIAQSVRIPQERKDR7IDFD	NDEDIKGILAAAKRAD	NIEDMQWGKGVREIPASVC	ik@lldtpnsravvi	DPPNIIIDYDEHKTM	CANGDPPIFTKPDKIS	CPRMSTIDGKELLGYIRA
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914	883	884	885	988	887	888	889	903	904	905	906	400	917	918	921	2693	2694	922	923
		-	•						,	•									
Q14833	P41594	P41594	P41594	P41594	P41594	P41594	P41594	015303	015303	015303	015303	015303	Q14831	Q14831	Q14831	Q14831	Q14831	000222	000222
Metabotropic Glutamate	Neceptor 4 Metabotropic Glutamate Receptor 5	Metabotropic Glutamate	Metabotropic Glutamate	Metabotropic Glutamate	Receptor 5 Metabotropic Glutamate	Metabotropic Glutamate	Metabotropic Giutamate Pecentor 5	Metabotropic Glutamate Receptor 7	Metabotropic Glutamate	Metabotropic Glutamate	Metabotropic Glutamate	Metabotropic Giutamate							
3006	3097	3097	3097	3097	3097	3097	3097	3098	3098	3098	3098	3098	3099	3099	3099	3066	3099	3100	3100
901	701	90	109	110	Ξ	112	113	114	. 311	911	117	118	119	120	121	122	123	124	125

	Homo sapiens	Homo saplens	Homo sapiens	Homo saniens	Homo sapiens	Homo sciplens	Homo sapiens	Homo sapiens	Homo sapiens		Homo saplens	-	Homo sapiens	Homo sapiens		Homo sapiens		Homo saplens	Homo sapiens		Homo saplens	Homo sapiens	1	Homo sapiens	Homo sapiens		Homo saplens	· .	Homo sapiens
	KVEDMQWAHREHTHPASVC	CESLETNISSIKTTYISYS	KFYWILTMMQRTHSQEYAHS	DGNISDPCGPNRTNLGGRDS	DRINHQLENLEAETAPLP	IKALVTIPETTFQTVS	RIRGNTRDHPSTANTVDR	SERSQPGAEGSPETPPGRC	CRAPRILQAYSWKEEE		SSEGEEPGSEVVIKMP		KQPPRSSPNTVKRPTKKGRD	CRWDKRRWRKIPKRPGS		EHNKIQNGKAPRDPVTENC		DSTSVSAVASNMRDDE	ENTVSTSLGHSKDENSKQTC		DEKONIVARKIVKMTK	RIKKDKKEPVANQDPVSPSL	• # * * * * * * * * * * * * * * * * * *	SKOKVHKHKPEGPKEKKAKI	KKPRPGGRPGGLRNGKLEEA		DKDTSNESSSGSATQNTKER		RPAANVARKFASIARNQVRK
	924	925	1894	231	232	233	234	1325	1326		1327		1328	1329		1330		1331	1332		1333	1831	910	017	219		220		5 2
	000222	000222	000222	AAA20580.1	AAA20580.1	AAA20580,1	AAA20580.1	AAA35686.1	AAA35686.1		AAA35686.1		AAA35686.1	AAA35686.1	,	AAA51570.1		AAA51570.1	AAA51570.1		AAA515/0.1	AAA51570.1	000615711	11701000	AAA51571.1		AAA51571.1		AAA51571.1
Receptor 8	Metabotropic Glutamate Receptor 8	Metabotropic Glutamate	Neceptor 8 Metabotropic Glutamate Receptor 8	Opioid mu-type Receptor	Opioid mu-type Receptor	Opioid mu-type Receptor	Opioid mu-type Receptor	Muscarinic acetylcholine Recentor M1	Muscarinic acetylcholine	Receptor M1	Muscarinic acetylcholine	Receptor MI	Muscarinic acetylcholine	Muscarinic acetylcholine	Receptor M1	Muscarinic acetylcholine	Receptor M2	Muscarinic acetylcholine Recentor M2	Muscarinic acetylcholine	Receptor M2	Muscarinic aceryicholine Receptor M2	Muscarinic acetylcholine	Receptor M2	Receptor M4	Muscarinic acetylcholine	Receptor M4	Muscarinic acetylcholine	Receptor M4	Muscarinic acetylcholine Receptor M4
	3100	3100	3100 .	3212	3212	3212	3212	3223	. 3223		3223	C	3223	3223		3224		3224	3224	7000	3224	3224	3006	277	3226		3226	,000	3220
	1126	1127	1128	1129	1130	1131	1132	1133	1134		1135	7611	9 -	1137		1138		6 13 6	1140	. נאונ	4.	1142	1143	<u>}</u>	1144		1145		40

Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens		Homo sapiens	Homo soniens	Homo sapiens	Homo sapiens	Homo sopiens	Homo soniens	Homo soniens	Homo sopiens	Homo sapiens		Homo sapiens	Homo saplens		Homo saplens	Homo sapiens	Homos caplens		Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
KAEKRKPAHRALFRSC	CSSYPSSEDEDKPATD	KESPGEEFSAEETEETFV	KFRLVVKADGNQETNNGC	KEPSTKGLNPNPSHQM		PAAEIWIDGGGGVGAD PSOPWANI TNOFVODSWP	SRKKRATPROPSENGC	ADAVNLTASLAAGAA	SPSALGLPVASPAPSQP	ERDFLPASDGTTTELVIRC	KTUKSAHNLPGEYNE	SEVARISSLDNSSFTAC	CGRKSYQERGTSYLLSSSA	RGELVPDPEPELIDST		CIVYHLESKISKRISF	REYSLIEIIPDFEIVAC		NDHYHQRRQKTTKMLVC	CEQRLDAIHSEVSVTFKAKK	MGPIGAEADENOTVEFMKVF		SEVSVTFKAKKNLEVRKNSG.	CVTVRQKEKANVTNLL	KNHSKALEFLADKVVC	CYARIYRRLQRQGRVFHKG
1334	1335	1336	1337	1338	1767	1759	1760	2265	2290	824	825	826	828	1057		1058	1059		1060	1061	2297		2298	1068	1069	1070
2	2	2	2		NP 001050 1	NP 001050.1	NP_001050.1	NP_001050.1	NP_001050.1		Ŷ	9	•			Q	•		9	9	• •		9			
P08912	P08912	P08912	P08912	P08912	<u>a</u> 2	3 0 2 2	N N	N 0.0	NP_0	P28336	P28336	P28336	P28336	P49146		P49146	P49146		P49146	P49146	P49146		P49146	P50391	P50391	P50391
Muscarinic Acetylcholine	Muscarinic Acetylcholine Recentor M5	Muscarlnic Acetylcholine	Muscarlnic Acetylcholine	Receptor M5 Muscarinic Acetylcholine	Receptor M5 Tochykinin Receptor 3	Tachykinin Receptor 3		Tachykinin Receptor 3		Neuromedin B Receptor	Neuromedin B Receptor	Neuromedin B Receptor	Neuromedin B Receptor	Neuropeptide Y Receptor	Type 2	Neuropeptide Y Receptor Type 2	Neuropeptide Y Receptor	z edyl	Neuropeptide Y Receptor Type 2	Neuropeptide Y Receptor	1ype 2 Neuropeptide Y Receptor	Type 2	Neuropeptide Y Receptor Ivoe 2	Neuropeptide Y Receptor Type 4	Neuropeptide Y Receptor	Neuropeptide Y Receptor
3227	3227	3227	3227	3227	3378	3378	3378	3378	3378	3380	3380	3380	3380	3404	Ç	3404	3404	,	3404	3404	3404		3404	3405	3405	3405
1147	1148	1149	1150	1151	1152	1153	152	1155		1157	1158	1159	1160	1161		701	1163 :	1144	<u> </u>	1165	1166	:	1167	1168	1169	1170

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	and sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens
TS IG INSSERI DE CE		SEHCQDSVDVMVFIVTS	MKKRNQKTTVNFLIGN	CGLSNKENRLEENEMI	NLTLHPSKKSGPQVKL	SFIKKHRRRYSKKTAC	PERPSQENHSRILPEN	CFEIKPEENSDVHELRV	RVLAAPSSELDVNTDIYS	CHPFKAKTLMSRSRTKK	GEQNRSADGQHAGGLVC	RQAAEQGQVCTVGGEHS	CPVWRRRRRPAFSRKADS	CHPIRALDVRTSSKAQA	PVAIMGSAQVEDEEIEC	GVQPSSETAVAILRFC •	CASALRRDVQVSDRVRSIAK	TPEPRPRTQPMASPRLGTFC	TAVASLLKGRQGIYTE
ובטו	2	2275	1072	1073	1074	1075	1076	1077	935	936	937	938	939	940	941	942	943	2123	2124
P50301		P50391	Q15761	Q15761	Q15761	Q15761	Q15761	Q15761	P30989	P30989	P30989	P30989	Туре Р30989	P41146	P41146	P41146	P41146	NP_000264.1	NP_000264.1
lype 4 Neuropeptide Y Receptor	Туре 4	Neuropeptide Y Receptor Type 4	Neuropeptide Y Receptor Type 5	Neuropeptide Y Receptor	Neurotensin Receptor Type	Oplate Receptor-Like 1	Oplate Receptor-Like 1	Oplate Receptor-Like 1	Opiate Receptor-Like 1	Ocular Albinism 1	Ocular Albinism 1 (Netfleship-Falls) (OA1)								
3405		3405	3406	3406	3406	3406	3406	3406	3408	3408	3408	3408	3408	3452	3452	3452	3452	3513	3513
1171		7/11	1173	1174	1175	1176	1177	1178	1179	1180	1181	1182	1183	1184	1185	1186	1187	1188	1189

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Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens		Homo sapiens	Homo sapiens		si iaidos oli ion	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens
EMQTDINGGSLKPVRTAAK	CSLGFQSPRKEIQWES	SEGSDASTIEIHTASESC	NPASGKVSQVGGQTSD	CKKLHIPLKAQNDLDISRIK	KIVKPLWTSFIQSVSYSKLL	TAITKKIFKSHLKSSRNSTS	VKKKSSRNIFSIVFVFFVC	AEGNRTAGPPRRNEALARVE	RLAVLATWLGCLVASAP	PEGAAAGDGGRVALAR	YLKGRRLGETSASKKSNSSS	MQRIGDVLGSSEDFRR		ARGGRATCHDISAPEL	KPAYGTSGGLPRAKRK		ופרטראור אונאמרפרואאטר	RYSGVVYPLKSLGRLKKKN	SGTGVRKNKTITCYD	RALIYKDLDNSPLRRKS	DIFRRISRATRKASRRSE	FV@STHSQGNNASEAC •	MVLKTLTKPVTLSRSKI	TIQNSIKMKNWSVRRSD	SEVHGAENFIGHNLQTLK	CTSRRALTRTAVYTLN	AGERRGKAARMAVVV
2125	2126	2127	2128	1486	1500	1502	1503	244	245	246	247	854	i L	622	856	967	??	386	387	388	389	850	851	852	853	874	875
																	-									÷	
NP_000264.1	NP_000264.1	NP_000264.1	NP_000264.1	NP_055694.1	NP_055694.1	NP_055694.1	NP_055694.1	CAA46097.1	CAA46097.1	CAA46097.1	CAA46097.1	AAC04923.1	. 000000	AACU4923.1	AAC04923.1	L 80000000	04423.1	CAA07339.1	CAA07339.1	CAA07339.1	CAA07339.1	P43657	P43657	P43657	P43657	Q15077	Q15077
Ocular Albinism 1	\mathcal{O}) <u> </u>	Ocular Albinism 1	(Nemesnip-ralls) (UAT) UDP-glucose Receptor	UDP-glucose Receptor	UDP-glucose Receptor (KIAA0001)	UDP-glucose Receptor (KIAA0001)	Oxytocin Receptor	Oxytocin Receptor	Oxytocin Receptor	Oxytocin Receptor			Pullnergic Keceptor PZY, G- protoin counted 2 (ppb/2)	Purinergic Receptor P2Y, G-	protein coupled, 2 (P2RY2)	protein coupled, 2 (P2RY2)		Purinergic Receptor P2Y1	Purinergic Receptor P2Y1	Purinergic Receptor P2Y1	_	Purinergic Receptor P2Y5	Purinergic Receptor P2Y5	Purinergic Receptor P2Y5	-	Purinergic Receptor P2Y6
3513	3513	3513	3513	3544	3544	3544	3544	3582	3582	3582	3582	3289	.000	2002	3589	3580		3595	3595	3595	3595	3596	3596	326	3596	,3297	3597
1190	1191	1192	1193	1194	1195	1196	1197	1198	1199	1200	120	1202	600	3	1204	1205	3.	1206	1207	1208	1209	1210	1211	1212	1213	1214	1215

																39	0/4	48																		
Homo saplens	Homo sapiens	Homo sapiens		Homo sapiens		Homo sabiens	-	Homo sapiens		Homo sapiens		Homo saplens	•	Homo sapiens		Homo sapiens		Homo sapiens		Homo sapiens		Homo sapiens		Homo sapiens	-	Homo sapiens		Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
TKTAYLAVRSTPGVPC	CHPI APWHKRGGDDAAW	CFRMKMRSETAIFITN		RTLRKPATLSQIGTNKK		ESFOKSFYINAHIRMES		KTETPLITKPSLPAIQEE		SSLRPRLGNATANNTCIVD		KAKVQCELNITAQLQEGE		ESLIMODDPQNSIEATSVDK		NSEQDCLPHSFHEETKE		EETKEDSGRQGDDILMEKPS		CEKRLKEVLQRPASIMESDK		ESEEDKEAPTGSRYRGRPC		LYSGATLDEAERLTEEELR		KDDGFLNGSCSGLDEEASG		CLEKIQRANELMGFNDSS	CPELFRIFNPDQVWETET	DSNSLDLSDMGVVSRNC	IKRKWRSWKVNRYFAVD	ESDFGDSNSLDLSDMGVVSR	RITGDLENTIKVQC	RSSREKRRSADIFIAS	GTIAGHFRKERIEGLRKRRR	GPNMGKGGEQMHEKSIPYSQ
876	2726	870		871		872	:	873		1895		248	-	249		250		251		761		762		763	•	765		944	945	946	948	2292	62	જ	28	92
Q15077	Q15077	Q99677		Q99677		G99677		G99677		G99677		AAC50157.1		AAC50157.1		AAC50157.1		AAC50157.1		Q03431		Q03431		Q03431	•-	Q03431	٠	P41586	P41586	P41586	P41586	P41586	AAA18954.1	AAA18954.1	AAA18954.1	AAA18954.1
Purinergic Receptor P2Y6	Purinergic Receptor P2Y6	G Protein-Coupled	Receptor 23 (GPR23)	G Protein-Coupled	Receptor 23 (GPR23)	G Protein-Coupled	Receptor 23 (GPR23)	G Protein-Coupled	Receptor 23 (GPR23)	G Protein-Coupled	Receptor 23 (GPR23)	Parathyrold Hormone	Receptor 2 (PTHR2)	Parathyroid Hormone	Receptor 2 (PTHR2)	Parathyroid Hormone	Receptor 2 (PTHR2)	Parathyroid Hormone	Receptor 2 (PTHR2)	Parathyroid Hormone	Receptor 1 (PTHR1)	Parathyroid Hormone	Receptor 1 (PTHR1)	Parathyroid Hormone	Receptor 1 (PTHR1)	Parathyroid Hormone	Receptor I (PIHRI)	PACAP Receptor Type 1	Apelin Receptor	Apelin Receptor	Apelin Receptor	Apelin Receptor				
3597	3597	3599		3266		3599	-	3599		3599		3638		3638		3638		3638		3640		3640		3640		3640		3732	3732	3732	3732	3732	3844	3844	3844	3844
1216	1218	1219		1220		1221		1222		1223		1224	٠	1225		1226		1227		1228		1229		1230		1231		1232	1233	1234	1235	1236	1237	1238	1239	1240

												3	911	/44	0														
Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens		Homo sapiens	Homo sapiens	Homo saplens		Homo saplens	-	• Homo sapiens	adolars omon		Homo saplens	Homo saplens
RMEDEDYNTSISYGDEYPD	DSIVVLEDLSPLEARVTR	LTIVCKLHRNRLAKTKKPFK	RSFTKMSSMNERTSMNERE	TRSRRLTFRKNISKASRSSE	CPSGDSAGKFKRPIIAG	CPSGDSAGKFKRPIIAGME	RSKSDNSSHPQKDEGD	ERHLTMIKMRPYDANK	LVKSSSRKVANHNNSE	SPKVKEDLPHTDPSSC	CLVRGRGARASPIQPALD	REHYQYVGKLAGRLKEASE	RAHTWREKRLLYSKMVC	KEESGIAICTMVYPSDEST	QAKKSSKHKALKVTIT	GERFRRDLVKTLKNLGC	ENYSYDLDYYSLESDLEEK		RDTVEFNNHTLCYNNFØKHD	SKKFQARFRSSVAEILK	GTVSEQLRNSETKNLC		HPLRRRISLRLSAYAV		CEEFWGSQERQRQLYA	COd/V (divd i//\s/\d/\s		CVTGSQADWDRARRRR	DSFREELRKLLVAWPRKIA
						<i>:</i>																							
447	844	449	450	1010	101	1012	1013	1028	1029	080	1031	1752	958	626	096	961	74		25	9/2	11		1087		1088	1080	2	1090	1001
		•					:																						
UR39	Q99788	Q99788	Q99788	AAA52336.1	AAA52336.1	AAA52336.1	AAA52336.1	Q99500	Q99500	Q99500	Q99500	Q99500	P51686	P51686	P51686	P51686	AAA64592.1		AAA64592.1	AAA64592.1	AAA64592.1		075194		075194	075104	5	075194	075194
Chemokine-Like Receptor 1	Chemokine-Like Receptor 1 (CMKLR1)	Chemokine-Like Receptor 1 (CMKLR1)	Chemokine-Like Receptor 1	Sphingolipid Receptor Edg1	Sphingolipid Receptor Edg1	Sphingolipid Receptor Edg1	Sphingolipid Receptor Edg1	Sphingolipid Receptor Edg3	C-C Chemokine Receptor 9	G Protein-Coupled	Receptor GPR1	G Protein-Coupled Receptor GPR1	G Protein-Coupled	Receptor GPR1 G Protein-Coupled	Receptor GPR1	G Protein-Coupled	Receptor 10 (GPR10)	G Protein-Coupled	(Protein-Coupled	Receptor 10 (GPR10)	G Protein-Coupled	Receptor 10 (GPR10) G Protein-Coupled							
3845	3845	3845	3845	3846	3846	3846	3846	3847	3847	3847	3847	3847	3848	3848	3848	3848	3849		3849	3849	3849		3850		3850	3850	3	3850	3850
241	242	243	244	245	246	247	248	249	220	251	252	253	254	255	256	257	258		259	260	261		262		263	26.4	3	265	266

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	Homo sapiens	Homo sapiens		Homo sapiens	Homo sapiens		Homo sapiens		Homo saplens	-	Homo saplens		Homo sapiens	•	Homo saplens		Homo saplens		Homo sapiens	•	Homo sapiens		Homo sapiens		Homo sapiens		Homo saplens	-	Homo saplens		Homo saplens		Homo saplens		Homo saplens	
	GCIPSSLAQRARSPSD	ENISAAVSSRVPAVEPEPE	* * T. * * T. * C. * * C. * C. * C. * C.	SICSVVIRPLIKININAA	QSEATKLVTIGUVAS		KOKENECLGDYPEVLOE		SMNNRTVQHGVTISL		ETLKLYDFFPSCDMRKDLR		GRSVHVDFSSSESQRSRHGS		CLKNYDFGSSTETSDSHLTK		KALSTFIHAEDFARRRRRS		ATSPNSDIRETHSHVP		LMGALHFKPGSRRUD		GLPTLLSRELTUDDKPYC		DRYMAIVQPKYAKELKNTC		KDPDKDSTPATCLKISD		GRTSKLKPKVKEKSIR •		RNYLRSLRRKSFRSGSLR	•	KVSREKAKKMIAASWIFD		DGRTVRRTMINIVPRTKVK	
	78	. 62			308		84		82		86		87		1511		1512		1612		1613		1615		93		24	79	95		%		26		86	
	AAA91630.1	AAA91630.1	. 007.0000	1.000.1	AAA91630.1		AAA91783.1	-	AAA91783.1	-	AAA91783.1		AAA91783.1		NP_005281.1		NP_005281.1		NP_005281.1		NP_005281.1	· · · · · · · · · · · · · · · · · · ·	NP_005281.1		AAB65819.1		AAB65819.1		AAB65819.1		AAB65819.1		AAB00316.1		AAB00316.1	
Receptor 10 (GPR10)	G Protein-Coupled Recentor GPR12	G Protein-Coupled	Receptor GPR12	Receptor GPR12	G Protein-Coupled	Receptor GPR12	CX3C Chemokine	Fractalkine Receptor 1	CX3C Chemokine	Fractalkine Receptor 1	CX3C Chemokine	Fractalkine Receptor 1	CX3C Chemokine	Fractalkine Receptor 1	G Protein-Coupled	Receptor GPR15	G Protein-Coupled	Receptor GPR15	G Protein-Coupled	Receptor GPR15	G Protein-Coupled	Receptor GPR15	G Protein-Coupled	Receptor GPR15	G Protein-Coupled	Receptor GPR18	G Protein-Coupled	Receptor GPR18	G Protein-Coupled	Receptor GPR18	G Protein-Coupled	Receptor GPR18	G Protein-Coupled	Receptor GPR19	G Protein-Coupled Receptor GPR19	····)
	3851	3851	205	200	3851		3852		3852		3852		3852		3853		3853		3853		3853		3853		3854		3854		3854		3854		3855		3855	
	1267	1268	10,00	607	1270		1271		1272		1273		1274		1275		1276		1277		1278		1279		1280		178		1282		1283		1284		1285	

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Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sanjens	Homo sonolens			Homo sapiens	Homo sapiens	Homo sapiens		Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homos carolens		Homo sapiens	Homo sapiens	Homo socions	a de la comon	2 22 22 20 20 20 20 20 20 20 20 20 20 20
RRGMKETFCMSSMKC	KTITKDSIYDSFDREAKEKK	ALLFSQDGQREGQRRC	SGDEEDAYSAEPLPELC	ALLIDTADILAARERSC	RRURGGSSPSGPQPRRGC	KGSGRHHII.SAGPHALTQ	DAINAS EL EN EULEADI DE		SKFGLLH&GRARKVKAIMQ	GQHGEREPSSGDVVSMHRSS	SERGARFSSGSGETGEVQAC		DPYTVRSKGPLNGC	NSTLDGNQSSHPFCLL	CASOTTANDPYTVRSK	EINMGSESNITVRDDIDD	RRAVKPHRERREROKRVERM		TRQKFQKVLKSKMKKR.	DPKRNKKITFEDSEIREKR	CAPGGGRRWRIPGPAWVFG	FASI I PTG-PNASNTSDG-PDN	
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8	8	1152	1153	1154	1155	101	5	2 5	3	<u>5</u>	105		9	107	98	109	=======================================		112	113	1532	1533	
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AAB00316.1	AAB00316.1	P46092	P46092	P46092	P46092	AAC51302.1	AAC51302 1	A A CE1200 1	74601302.1	AAC51302.1	AAC51303.1		AAC51303.1	AAC51303.1	AAC51303.1	AAC51304.1	AAC51304.1		AAC51304.1	AAC51304.1	AAH01736.1	AAH01736.1	
G Protein-Coupled	G Protein-Coupled Receptor GPR19	G Protein-Coupled Receptor GPR2/CCR10	G Protein-Coupled Becontar CBD2/CCB10	G Protein-Coupled	Receptor GPR2/CCR10 G Protein-Coupled	Receptor GPR2/CCR10 G Protein-Coupled	Receptor GPR20 G Protein-Coupled	Receptor GPR20	Receptor GPR20	G Protein-Coupled	G Protein-Coupled	Receptor GPR21	G Protein-Coupled Receptor GPR21	G Protein-Coupled Receptor GPR21	G Protein-Coupled	G Protein-Coupled	Receptor GPR22 G Protein-Coupled	Receptor GPR22	G Protein-Coupled Pecentor GPD22	G Protein-Coupled	receptor GPR22 G Protein-Coupled	Receptor SLC/MCH1 © Protein-Coupled	
3855	3855	3856	3856	3856	3856	3857	3857	3857	3	3857	3858		3858	3858	3858	3859	3859		3859	3859	3860	3860	-
1286	1287	1288	1289	1290	1291	1292	1293	1204	1	1295	1296		1297	1298	1299	1300	1301		1302	1303	1304	1305	٠.

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	WC	02/0	06108	37 .		,			-	39	4/448	,	e.				PCT	r/US0	1/501	.07
	Homo sapiens	Homo saplens	Homo sapiens	•	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
	KGVGRAVGLGGGSGCQATE	RMTSSVAPASQRSIRLRTKR	RAVSNAQTADEERTESKG		RGLQPLPGGQDSQCGEEP	CRISRRLRRPPHVGRARRNS	RTGRLARRISSASSLSRDD	DYSGLDGLEELELCPAGD	TVYCLLGDAHSPPLYT	EGPTGPAAPLPSPKAWD	HFAAVFCIGSAEMSL	GLTCGVVYPLSKNH	REPEKQPKLQRAQALVTLV	CHSFYSRADGSFSIIWQEA	QNLGSCRALCAVAHTSDVTG	SPTFRSSYRRVFHTLRGKGQ	DELFRDRYNHTFCFEKFPME •	LRAVRGSVSTERQEKAKIKR	RSDVAKALHNLLRFLASDK	NASLTLETPLTSKRNSTAK
	1539	1565	1567		376	377	378	483	118	911	120	121	1157	1158	1159	1160	143	144	145	146
	AAH01736.1	AAH01736.1	AAH01736.1		3000	000155	000155	000155	AAB60402.1	AAB60402.1	AAB60402.1	AAB60402.1	000270	000270	000270	000270	AAA98457.1	AAA98457.1	AAA98457.1	AAA98457.1
Receptor SLC/MCH1	G Protein-Coupled	G Protein-Coupled	Receptor SLC/MCHI G Protein-Coupled	Receptor SLC/MCH1	G Protein-Coupled Receptor GPR25	G Protein-Coupled	G Protein-Coupled	Receptor GPK25 G Protein-Coupled	Receptor GPK25 G-Protein-Coupled	G Protein-Coupled	Keceptor GPK3 G Protein-Coupled	Keceptor GPKS G Protein-Coupled	Receptor GPRs G Protein-Coupled	G Protein-Coupled	Receptor Grivsi G Protein-Coupled	G Protein-Coupled	receptor GPR3 G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	receptor GPR4 G Protein-Coupled Receptor GPR4

1321 - 3864

Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
FQYLVPSETVSILTVG	CLAERAACSVVRPLARSH	HLYVRICQVVWRHAH	EIGRALWLLCGCFGSK	ATAESRRVAGRTYSAAR	RLDDEGGRRGCVLVFPQPE	RIHAMRIDSHAKALERAKKR	DASFRRNLRQUTC	NVSQDNGTGHNATFSEP	RSRHMPWRTYRGAKVAS	VRLRSGAKALGKARRK	LDDNFRKNFRSILRC	QDHFLEIDKKNCCVFRDD	ARIIWSLRQRQMDRHAKIKR	CLQRKMTGEPDNNRSTSVE	DPNKTRGAPEALMANSGE	SNNHSKKGHCHQEPASLEKQ	RGRQMDRHAKIKRAITFIMV	SPSYLGPTSNNHSKKG	AVRRSHGTQKSRKDQI
166	167	168	169	171	172	173	174	175	176	771	178	179	180	181	182	183	1453	1454	1192
AAA91631.1	AAA91631.1	AAA91631.1	AAA91631.1	AAC50197.1	AAC50197.1	AAC50197.1	AAC50197.1	AAC50198.1	AAC50198.1	AAC50198.1	AAC50198.1	BAA01721.1	BAA01721.1	BAA01721.1	BAA01721.1	BAA01721.1	BAA01721.1	BAA01721.1	Q15743
G Protein-Coupled Receptor GPR6	G Protein-Coupled Receptor GPR6	G Protein-Coupled Receptor GPR6	G Protein-Coupled Receptor GPR6	G Protein-Coupled Receptor GPR7	G Protein-Coupled Recentor GPR7	G Protein-Coupled Receptor GPR7	G Protein-Coupled Receptor GPR7	G Protein-Coupled Recentor GPR8	G Protein-Coupled Receptor GPR8	G Protein-Coupled	G Protein-Coupled Receptor GPR8	G Protein-Coupled	G Protein-Coupled Receptor HM74	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled Recentor HM74	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled
3866	3866	3866	3866	3867	3867	3867	3867	3868	3868	3868	3868	3869	3869	3869	3869	3869	3869	3869	3870
1325	1326	1327	1328	1329	1330	1331	1332	1333	1334	1335	1336	1337	1338	1339	1340	1341	1342	1343	1344

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	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens
-	LMHEEVIEDENQHRVC ·	CFVSETTHRDLARLRG	CSRTGRAREAYPLGAPEASG	CRMYRQQKRHQGSLGPRPRT	CFTQAVAPDSSSEMGD	ASGRRDPRAPSAPVGKEGSC	SAWGEGQVEPLPPTQQ	KSPFYRCQNTISVEKGNSAV	RNLYAMHRRLQRHPRSC	CAEPRADGREASPQPLEEL	KDVKEKNRTSEEAEDLRALR	AQAAGRLRRRSATTF	CVGVTRPLLHAARVSVARAR	CNILSGLALHRARWRR	ASGPDSRRRWGAHGPR	SGSARRARAHDVEMVGQ	IALALLARRWRGDVGC	CETRQWLPPGESPAISSV	GPSLGSGRGGPGARRRGE	NETSSRKEKWDLQ.ALR	ERSAEARGNLTRPPGSGEDC	SRSYRRESKRKKSFLLC	CRAKATASQSSAQWGR
	1193	1194	1195	1188	1189	1190	1191	458	459	503	504	962	963	964	996	996	196	896	696	971	972	973	974
	-																						
•	Q15743	Q15743	Q15743	P43119	P43119	P43119	P43119	Q13258	Q13258	Q13258	Q13258	P34995	P34995	P34995	P34995	P34995	AAD44177.1	AAD44177.1	AAD44177.1	AAD44177.1	CAB52459.1	CAB52459.1	CAB52459.1
Receptor OGR1	G Protein-Coupled Receptor OGR1	G Protein-Coupled Receptor OGR1	G Protein-Coupled Recentor OGR	Prostacyclin Receptor	Prostacyclin Receptor	Prostacyclin Receptor	Prostacyclin Receptor	Prostaglandin D2 Receptor	Prostaglandin D2 Receptor	Prostaglandin D2 Receptor	Prostaglandin D2 Receptor	Prostaglandin E Receptor EP1	Prostaglandin E Receptor EP1	Prostaglandin E Receptor	Prostaglandin E. Receptor FP1	Prostaglandin E Receptor 7	Prostaglandin E Receptor EP2	ptor	Prostaglandin E Receptor FP2	Prostaglandin E Receptor FP2	Prostaglandin E2 Receptor FP3	Prostaglandin E2 Receptor	Prostaglandin E2 Receptor
	3870	3870	3870	3921	3921	3921	3921	3923	3923	3923	3923	3924	3924	3924	3924	3924	3925	3925	3925	3925	3926	3926	3926
	1345	1346	1347	1348	1349	1350	1351	1352	1353	1354	1355	1356	1357	1358	1359	1360	1361	1362	1363	1364	1365	1366	1367

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	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	
	KFCQVANAVSSCSNDGQ	RLSDFRRRRSFRRIAGAE	EREVSKNPDLQAIRIAS	DSQRTSSAMSGHSRSFISRE	RTLRISETSDSSQGQDSE	ILMKAYQRFRQKSKAS	ASDKEWIRFDOSNVLC	TKPIFHSTKITSKHVK	CFYNTEDIKDWEDRFY	RVKFKSQQHRQGRSHHLE	QGTNRSSKGRSLIGKVDGTS	GRYWVIVNPMGHSRKKAN	SHDFRDHAKNALLCRSVR	VSLTSKKHSRKSSSYS	ENDTNNLAKPTLPIKTFR	CPEESASHLHVKNATMG	QPDITTCHDVHNTCESSSP	MSKTRNHSTAYLTK	RDHKSGTPANVFLMH	
-	975	382	383	384	385	1046	1047	1048	1049	1050	252	253	255	256	257	258	260	261	88	
	CAB52459.1	P35408	P35408	P35408	P35408	P43088	P43088	P43088	P43088	P43088	AAB47871.1	AAB47871.1	AAB47871.1	AAB47871.1	AAC51218.1	AAC51218.1	AAC51218.1	AAC51218.1	CAB08108.1	
. T	Prostaglandin E2 Receptor EP3	Prostaglandin E Receptor EP4	Prostaglandin E Receptor EP4	Prostaglandin E Receptor EP4	Prostaglandin E Receptor EP4	Prostaglandin F2-alpha Receptor	Prostaglandin F2-aipha Receptor	Prostaglandin F2-alpha	Prostaglandin F2-alpha Recentor	Prostaglandin F2-alpha Receptor	Proteinase-Activated Receptor 2	Proteinase-Activated Receptor 2	Proteinase-Activated Receptor 2	Proteinase-Activated Receptor 2	Proteinase-Activated Receptor 3	Proteinase-Activated Receptor 3	Proteinase-Activated Receptor 3	Proteinase-Activated	G Protein-Coupled Receptor GPR17	
	3926	3927	3927	3927	3927	3928	3928	3928	3928	3928	4051	4051	4051	4051	4052	4052	4052	4052	4090	
	368	369	370	371	372	373	374	375	376	377	378	379	380	381	382	383	384	385	386	

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Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	,	Homo sapiens	Homo sopiens		Homo sapiens		Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens
RSLRGGLRVEKRLKTKAVR	RSHGASCATQRILALANR	FEGKTNESSL\$AKSE	RNCMLTTICCGKNPLGD	CGIDYYTLKPEVNNESFVI	CWVPYASVAFYIFTHQGSN	VLGGFTSTLYTSLHGY	ATSSLLRRWPYGSDGC		CTLDYSKGDRNFTSFL	MEGKIGKSGHLOVNIT		MVCRGIWQCLSPQKRE		CLQELSREQTGDLGTEQ	CPRFLRMLTSRNGSLFRN	CGVNVNDSSNEKRHSY	KDAVLFSSDDVTYCDAH	MRKLRTGETRGNEVSH	EEPGRNASQNGTLSEG	CLSWMDNAAEEPVDY	EDFQPENLESGGVFRNGTC	LSVDAVNMFTSIYC	RAYSVEDFQPENLES	RSNQWGRSSCTINWPGE	KVKSSGIRVGSSKRKKSE	CLVKVSGTDDGERSDS
			051	1052	1053	1055	1042		1043	1044		1045		920	951	952	954	926	994	966	266	2616	2618	866	666	1000
06	16)[2	×		=		= .			=		8	8	ŏ	ŏ	ŏ	ŏ	ŏ		Š	, ở	Ŏ.	δ.	
CAB08108.1	CAB08108.1	CAB08108.1	P08100	P08100	P08100	P08100	P47804		P47804	P47804		P47804		P47872	P47872	P47872	P47872	P47872	P30872	P30872	P30872	P30872	P30872	P30874	P30874	P30874
G Protein-Coupled	Receptor GPR17 G Protein-Coupled Receptor GPR17	G Protein-Coupled Receptor GPR17	Rhodopsin	Rhodopsin	Rhodopsin	Rhodopsin	Retinal G Protein-Coupled	Receptor RPE	Retinal G Protein-Coupled:	Refinal G Protein-Coupled	Receptor RPE	Retinal G Protein-Coupled	Receptor RPE	Secretin Receptor	Secretin Receptor	Secretin Receptor	Secretin Receptor	Secretin Receptor	Somatostatin Receptor Type							
4090	4090	4090	4254	4254	4254	4254	4284		4284	4284		4284		4321	4321	4321	4321	4321	4480	4480	4480	4480	4480	4481	4481	4481
1387	388	389	1390	1391	392	393	394		395	396		397	٠.	398	336	40	6	402	403	404	405	406	1407	1408	1409	1410

	399/448		PCT/US01/50107
Homo saplens Homo saplens Homo saplens	Homo sapiens Homo sapiens Homo sapiens	Homo sapiens Homo sapiens Homo sapiens	Homo saplens Homo saplens Homo saplens Homo saplens Homo saplens Homo saplens
TISEPENASSAWPPD QPGTSGQERPPSRVA IFADTRPARGGQAVAC	CLLEGAGGAEEPLDY KMRAVALRAGWQQRR CRAVLSVDGLNMFTSV CLVGLVGNALVIFVIL	SLPLLVFADVQEGGTC CLRKGSGAKDADATEP RIRQQQEATPPAHRAAA RVAKLASAAAWVUSLC	CMIEWPEHPNKIYEKV CPFISAGDYEGLEMKSTRYL KVSRLETTISTVVGAHEE EPEDGPKATPSSLDLTSNC EDEEKNESGLTEYRLV AVANRSKKSRALFLSAAVFC SINKSSPLQKQLPAFISE
4	3 1 7 8	3 9 8	000000
262 262 100	100 262 263 263 263	263 263 264 264	1339 1340 1341 1202 1202 2583 2583
		144.1 144.1 145.1	41.1 41.1 1.1 1.1
e P32745 e P32745 e P31391	6 P31391 6 P31391 6 P31391 6 P31391	MP_001C	AAA36641. AAA36641. AAA36641. AAA36641. P25116 P25116
omatostatin Receptor Typ omatostatin Receptor Typ omatostatin Receptor Typ	amatostatin Receptor Typ amatostatin Receptor Typ amatostatin Receptor Typ amatostatin Receptor Typ	matostatin Receptor Typ matostatin Receptor Typ matostatin Receptor Typ matostatin Receptor Typ	5 Tachykinin Receptor 1 Tachykinin Receptor 1 Tachykinin Receptor 1 Tachykinin Receptor 1 Thrombin Receptor 1
	32745 2624 TTSEPENASSAWPPD 32745 2626 QPGTSGQERPPSRVA 31391 1007 IFADTRPARGGQAVAC	32745 2624 TTSEPENASSAWPPD Homo sapiens 32745 2626 QPGTSGQERPPSRVA Homo sapiens 331391 1007 IFADTRPARGGQAVAC Homo sapiens 331391 2627 KMIRAVALRAGWQQRR Homo sapiens 331391 2633 CLVGLVGNALVIFVIL Homo sapiens 33391 2633 CLVGLVGNALVIFVIL Homo sapiens	P32745 2624 TTSEPENASSAWPPD Homo sapiens P32745 2626 QPGTSGQERPPSRVA Homo sapiens P31391 1007 IFADTRPARGCQAVAC Homo sapiens P31391 1008 CLLEGAGGEEFPLDY Homo sapiens P31391 2627 KMRAVALRAGWQGRR Homo sapiens P31391 2637 KMRAVSVDGLNIMFTSV Homo sapiens P31391 2633 CLVGLVGNALVIFVIL Homo sapiens NP_001044.1 2637 SLPILVFADVQEGGTC Homo sapiens NP_001044.1 2639 RIRGQGEATPPAHRAAA Homo sapiens NP_001044.1 2643 RVAKLASAAAWVISLC Homo sapiens

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Homo sapiens			Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens
DPRSFLLRNPNDKYEPFWE	CENSTVSSRKGNTKMI A		KAATIKALCINCARKPIE	KPANYSVALNYSVIKE	KESDHFSTELDDITVTD	EIQKNKPRNDDIFKII	SYRPSDNVSSSTKKPAPC	LNSSTEDGIKRIQDDC	CSQKPSDKHLDAIPIL	DRYGSVIYPFLSQRRN	RKHLLKTNSYGKNRITRD	RVPITWLQGKRESMSC	CHDTTRPEEFDHYVHFSSA	YLLTGDKYRRQLRQLC	HPLRALRWGRPRLAG	HITRTIYYLARLLEADC •	REAEALGEGNGPPRDVRNEE	NVRGKTASRQSKGAEQ	QNMKEKFNKEDTDSMSRRQ	RQTFYSNNRSPTNSTGMWKD	NATTPWLGRDEELAKVE	TRGLPSRVSSINTISRAKIR
2621	1197		0 1	19%	1200	1771	1772	1773	1321	1322	1323	1324	1142	1145	2696	2697	262	263	264	265	266	267
P25116 P34981	P34981	D34081	104401	P34981	P34981	NP_000676.1	NP_000676.1	NP_000676.1	P50052	P50052	P50052	P50052	P51582	P51582	P51582	P51582	AAA62271.1	AAA62271.1	AAA62271.1	AAA62271.1	AAA65687.1	AAA65687.1
Thrombin Receptor Thyrotropin Releasing	Hormone Receptor Thyrotropin Releasing	Hormone Receptor Thyrotropia Delegating	Hormone Receptor	Thyrotropin Releasing Hormone Receptor	Thyrotropin Releasing Hormone Receptor	Anglotensin II Type 1 Receptor	Angiotensin II Type 1 Receptor	Angiotensin II Type 1 Receptor	Angiotensin II Type 2 Receptor	Pyrimidinergic Receptor P2Y4	Pyrimidinergic Receptor P2Y4	Pyrimidinergic Receptor P2Y4	Pyrimidinergic Receptor P2Y4	Vasopressin V1A Receptor	Vasopressin V1A Receptor	Vasopressin V1A Receptor	Vasopressin V1A Receptor	Vasopressin V1B Receptor	Vasopressin V1B Receptor			
4687	4734	4734	î Î	4734	4734	4944	4944	4944	4946	4946	4946	4946	5072	5072	5072	5072	5117	5117	2117	5117	5118	5118
1433	1435	1/34	3	1437	1438	1439	1440	1441	1442	1443	1444 4	1445	1446	1447	1448	1449	1450	1451	1452	1453	454	1455

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	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	مرواطي وسولا		Homo sapiens	Homo sapiens		Homo sapiens		Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homos cmcH		Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
	ESPRDLELADGEGTAET	SNSSGERPLDTRDPLLARAE	RHGSGAHWNRPVLVAWAFS	CQVLIFREIHASLVPGPSER	RGRIPPSLGPQDESC	KNEDGSVFSQTEHNIV	IKYKELRTPTNAIIIN	RKNDRSFVSYTMTVIA	CTESLNRDWSDQIDVTK	VANKKFRRAMLAMFKC	CGPAGRISSRSQSLRSTDAR	EENIDOMAKEEAO! ACBN		CRVVDRQEEGNGDSGG	KRDKAPKSSFVGDGDI		RKLQHAAEKDKEVLGP		CLRPSPEEAVAQAESEVGR	GSSNDLFTTEMRYGEE	MARDGISDKSKKQRAGSERC	EDAPRARPEGIPRRAAK	PSPTMPPT/PGSTMKNAGS1 F		KREKRWSVSSGGAAERSVC	RRVFPTNFPGLQKKGE	CNLTREAKRPPKEEFG	KLKHRAGQMSEPHSGLTLKC
348	269	270	271	272	273	1147	1148	1149	1150	1151	286	880	2	686	066	Projection of the control of the con	166		981	982	983	984	985	3	986	976	977	978
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0 0 0 65587 1	AAA65687.1	CAA77746.1	CAA77746.1	CAA77746.1	CAA77746.1	014718	014718	014718	014718	014718	014514	014514	<u>;</u>	014514	014514		014514	.,	060241	060241	060241	060241	O60241	Ţ	O60241	O60242	060242	060242
Vasoprassip VIB Deceptor	Vasopressin V18 Receptor	Vasopressin V2 Receptor	Vasopressin V2 Receptor	Vasopressin V2 Receptor	Vasopressin V2 Receptor	Peropsin	Peropsin	Peropsin	Peropsin	Peropsin	Brain-Specific Anglogenesis	Innibilor 1 Brain-Specific Apalogenesis	Inhibitor 1	Brain-Specific Anglogenesis	Brain-Specific Anglogenesis	Inhibitor 1	Brain-Specific Anglogenesis	٠.		Brain-Specific Angiogenesis Inhibitor 2	Brain-Specific Angiogenesis Inhibitor 2	Brain-Specific Anglogenesis	Innibitor 2 Brain-Specific Anaiogenesis	Inhibitor 2	Brain-Specific Anglogenesis Inhibitor 2		Brain-Specific Angiogenesis Inhibitor 3	Brain-Specific Angiogenesis
5118	5118	5119	5119	5119	5119	5133	5133	5133	5133	5133	5519	5519	}	5519	5519		5519		92ZQ	5520	5520	5520	5520		5520	5521	5521	5521
1454	1457	1458	1459	9460	1461	1462	1463	1464	1465	1466	1467	1468) }.	1469	1470		1471	47	14/2	1473	1474	1475	1476) · :	1477	1478	1479	1480

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Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens		Homo saplens	Homo saplens		Homo sapiens		Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens		Homo sapiens	Homos omor		Homo sapiens		Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens
CIDDNLRGADMDIVHPGER	SRSETGSTISMSSLERR	NDSSQEEHQDFLQFSK	KATKAYNQQAKRMTWG	KTLLHAGGFQKHRSLK	SLKFRKNFWKLVKDIGC	KSSEDNSKTFSASHNV	ERHRSVMAVQLHSRLPRGR		RRRVQRMAEHVSCHPRYRE	NAAVYSCRDAEMRRIFRR		ROSTRESVHYTSSAQGGAST		YSQYQFWKNFQTLK	QQEAPERASSVYTRSTGEQE	RSQKEGLHYTCSSHFPYSQ	MDYQVSSPIYDINYYTSEPC	EDEYDVLIEGELESDEAEQC		KGNFFSARRRVPCGIITSVL	MAPKTI DEDECIDASI EKI VEA		RSNTPLQPRGQSAQGTSRE		GPGNSARDVLRARAPREEQG .	DPGGPRRGNSTNRRVRLKNP	LRQLSKEDLGFSGRAPAERC	PRGAVISGRSQEQSVKTVPG	CIQKSSTVTSDDNDNEYTTE	CIQKSSTVTSDDNDNEYTTE	TDVVETRLSQWLEEMPC
676	086	1101	1102	1103	1104	1105	%		29	. 89		69		38	39	40	309	1092		1093	1004	<u>.</u>	9601		127	129	130	131	1781	1806	319
060242	O60242	000574	000574	000574	000574	000574	AAC27728.1		AAC27728.1	AAC27728.1		AAC27728.1		AAC50598.1	AAC50598.1	AAC50598.1	AAC50598.1	O00421		000421	000421	(((1)	000421		AAC51281.1	AAC51281.1	AAC51281.1	AAC51281.1	AAC51281.1	NP_005293.1	014804
Inhibitor 3 Brain-Specific Angiogenesis Inhibitor 3	cific Angiogenesis	SIV/HIV Receptor BONZO	SIV/HIV Receptor BONZO	SIV/HIV Receptor BONZO			Lysophosphatidic Acid	٠.	Lysophosphatidic Acid	Jic Acid		Lysophosphatidic Acid	Receptor Edg4	Chemokine Receptor 5	C-C Chemokine Receptor 5	C-C Chemokine Receptor 5	C-C Chemokine Receptor 5	Chemokine (C-C motif)	Receptor-like 2 (CCRL2)	Chemokine (C-C motif)	~ <i>-</i>	.	Chemokine (C-C motif)	Receptor-like 2 (CCRL2)	Pael Receptor (GPR37)	Pael Receptor (GPR37)	Pael Receptor (GPR37)	Pael Receptor (GPR37)	Pael Receptor (GPR37)	Pael Receptor (GPR37)	Putative Neurotransmitter Receptor (PNR)
5521	5521	6031	.6031	6031	6031	6031	6204	,	6204	6204		6204		6213	6213	6213	6213	6363	:	6363	6363	}	6363		6446	6446	644 6	6446 6446	6446	6446	6536
481	482	483	484	485	486	487	488	9	489	490	. •	491		492	493	494	495	496	!	497	498) : :	499		8	<u>8</u>	205	503	1504	1505	206

	485 HPAAFCYQVNGSCPR 788 KAKSKYSPELLKYRLP	788 KAKSKYSPELLKYRLP		290 KTGNWERKVIVSVRVA Homo saplens	60478 T91 KSVHSFDYDWYNVSDQAD Homo saplens	360478 TOP2 RVRNPTKDLTNPGMVP Homo saplens	060478 793 RYDSDDDLAWNIAPQGLQ Homo sapiens	43190 865 PTLSFSHLKRPQQGAGNC Homo sapiens	866 GALGRAVLRSPGMTVAE	867 MRVLNVDARRRWSTRC	-	2299	2300		AC26082.1 Homo sapiens	AC26082.1 FKRLRVHAHSTTDSAR Homo sapiens	AC26082 1 141 VORPHIFASPROSSARPTEK Homo soniens		AC26082.1 142 QSEAEPQSKSQSLSLESLEP Homo saplens	AC39634.1 • Homo sapiens	1 RAVDPVAAGSGARRAKRK	199 GRAPGRASGRVCAAARG	.1 200 ERESSDLLHMSEAAGALRPC	_	AC39601.1 236 EPSATPGAQMGVPPGSR Homo saplens	
		HPAAFCY(KAKSKYSPI	KTGNWER	KSVHSFDY	RVRNPTKC	RYDSDDDI	PTLSFSHLKI	GALGRAV	MRVLNVD	CPGYRDS	CPANFLA	ASNGLAL	CNRSSTRH	PNQIRRIM	EKRLRVHA	VORPHIFA		QSEAEPQ	NLTVCHP	RAVDPVA	GRAPGRA	ERESSDLL	DOLGDLE	EPSATPG.A	
		485	788	790	791	792	793	865	998	. 498	868	2299	2300	137	139	140	141		142	197	198	199	200	235	236	
		014804	O60478	060478	060478	060478	060478	043190	043190	043190	043190	043190	043190	AAC26082.	AAC26082.	AAC26082.	AAC26082		AAC26082.	AAC39634.	AAC39634.	AAC39634.	AAC39634.	AAC39601.	AAC39601.	
	Receptor (PNR)	Purative Neurotransmitter Receptor (PNR)	G Protein-Coupled Receptor TM7SF1	G Protein-Coupled Receptor TM7SF1	G Protein-Coupled Receptor TM7SF1	G Protein-Coupled Receptor TM7SF1	G Protein-Coupled Receptor TM7SF1	Purinergic Receptor P2Y11	Purinergic Receptor P2Y11	Purinergic Receptor P2Y11	Purinergic Receptor P2Y11	Purinergic Receptor P2Y11	Purinergic Receptor P2Y11	G Protein-Coupled Receptor GPR39	G Protein-Coupled	G Protein-Coupled	receptor GPR39 G Protein-Coupled	Receptor GPR39	G Protein-Coupled Receptor GPR39	Galanin Receptor GalR2	Galanin Receptor GalR2	Galanin Receptor GaiR2	Galanin Receptor GalR2	Orexin Receptor 1	Orexin Receptor 1	
6536	Č	000	7779	.7779	2777	7779	7779	6853	6853	6853	6853	6853	6853	6921	6921	6921	6921	· ·	6921	7221	7221	7221	7221	7246	7246	
1508		25	1510	1511	1512	1513	1514	1515	1516	1517	1518	1519	1520	1521	1522	1523	1524		1525	1526	1527	1528	1529	1530	153	

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Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens		Homo sapiens		Homo sapiens		Homo sapiens		Homo sapiens		Homo saplens		Homo saplens		Homo saplens	•	Homo saplens		Homo saplens		Homo sapiens	11	uoino sapiens	acias, cmcl	supidos oution	Homo saplens	· · · · · · · · · · · · · · · · · · ·	Homo sapiens		Homo saplens	Homo saplens	
KRPSDQLGDLEQGLSGEPQ	SELNET © EPFLNPT DYDDEE	KWKPLQPVSQPRGPGQ	TKSRMSAVAAEIKQIRA	RGEDRLTRGRTSTESRKS	AVTRPIKTAQANTRKR	· · · · · · · · · · · · · · · · · · ·	DSTNTVPDSAGSGNVTRC		QQRNAEVKRRALWMVC		KKFRKHLTEKFYSMRSSRKC		DRYYSVLYPLERKISDAKSR		DEEESEAKYIGSADFQAKE		ETRNSKKRLLPPLGNTPEE		ELIQTKVPKVGRVERKMSR		KKQRKAQNFTSILIAN		FRNLSLPTDLYTHQVAC		CVENWPSKKDRLLFTT		CLAKKANANVONNNEINEGIK	DEDECONIVII DAVIDIVA		CYFKIYIRLKRRNNMMDK		CDFRSRDDDYETIAMS		ENDUCHLPLAMIFILALA	SNFSEKNAQLLAFENDDC	
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237	240	241	245	243	1097		1098		1099		138	٠.	398		40	-	4		402		1078		1079		1080	1001	<u> </u>	1061	5	1065		1066		1498	2291	
															-											.•										
AAC39601.1	AAC39602.1	AAC39602.1	AAC39602.1	AAC39602.1	P25105	÷.	P25105		P25105		P25105		Q14439		Q14439	• .	Q14439		Q14439		Q99463		Q99463	:	Q99463	000463	644400	P05000	(7)(7)	P25929		P25929	000	F25929	P25929	
Orexin Receptor 1 Orexin Receptor 1	Orexin Receptor 2	Orexin Receptor 2	Orexin Receptor 2	Orexin Receptor 2	Platelet-Activating Factor	Receptor	Platelet-Activating Factor	Receptor	Platelet-Activating Factor	кесертог	Platelet-Activating Factor	Receptor	G Protein-Coupled	Receptor L8509	G Protein-Coupled	Receptor Ls8509	G Protein-Coupled	Receptor L8509	G Protein-Coupled	Receptor L88509	Neuropeptide Y Receptor	Type 6 Pseudogene	Neuropeptide Y Receptor	Type 6 Pseudogene	Neuropeptide Y Receptor	lype o Pseudogene	Type 4 Peer dogene	Neuropentide V Receptor		Neuropeptide Y Receptor	Type 1	Neuropeptide Y Receptor	lype l	Neuropeptide Y Receptor Type 1	Neuropeptide Y Receptor	
7246	7247	7247	7247	7247	8436	٠,	8436		8436		8436		8203		8509		8209		8209		988		8896		9886	8088	8	1670		9421		9421		7471	9421	
1532	1534	1535	1536	1537	1538		1539		1540		<u> </u>		1542	ř.	1543		154		1545		1546		1547		1548	15,40	<u>}</u>	1550	3	1551		1552	,,,,,	200	1554	

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	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens		Homo saplens		Homo sapiens		Homo sabiens	 		Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens
	CESLSLASNISDNGYRE	CGEILNEEKKSKVHYHVA	NHSEDGAPALLTTAPP	GGAPPRYATIEHPFHC	CEPARPDGSMFFSQEE	AAREAGAAVRRPLGPE		LRYRRPPREKIGRRRA		PRELAAGOSFHGCLYR		CKTVRLSDVRVRPVNTYAR			EDFWKGEDLSNYSYSS	PPFLLDAAPCEPESLE	RRTVYSSNVSPACYE	SKDSLPKDSRPSFVGS	PKPFLYVVGRKKMMDAQYKC	VEVVPNGELVRRDPVSC	KIQWNQRWGRRPSNRS	CHQEPRNEPANNQGEESAE	TKSFRLRSRTLPRSKIIC	STFVFNQKYNTQGSDVCE	TAANLGKMNRSCOSE	RYSENISRGTSETADNDNAS	CPLAPPELHPPAPAP	CAIVERERGWPDFLR	CINEVQNIKFNSSGQ	CEVPLVRTDNPKSWYE	CRADGTMRLGEPTSNE
	1778	1779	1774	1775	1776	1082		1083		1085		1086			802	803	804	805	992	. 692	177	772	355	356	357	358	2595	2666	2667	2668	599
	NP_004373.1	NP_004373.1	NP_001457.1	NP_001457.1	NP_001457.1	AAB97766.1		AAB97766.1		AAB97766.1		AAB97766.1			P25025	P25025	P25025	P25025	P30988	P30988	P30988	P30988	P51684	P51684	P51684	P51684	NP_005622.1	NP_005622.1	NP_005622.1	NP_005622.1	NP_005622.1
- DO	Corticotropin releasing factor Receptor 1	asing	Frizzled-2	Frizzled-2	Frizzled-2	Putative Leukocyte Platelet-	Activating Factor Receptor (HUMINPIIY20)	Putative Leukocyte Platelet-	Activating Factor Receptor (HUMNPIIY20)	ocyte Platelet-		Putative Leukocyte Platelet-	Activating Factor Receptor	(HUMINPIIY20)	Interleukin-8 Receptor B	Interleukin-8 Receptor B	Interleukin-8 Receptor B	Interleukin-8 Receptor B	Calcitonin Receptor	Calcitonin Receptor	Calcitonin Receptor	Calcitonin Receptor	C-C Chemokine Receptor 6	Smoothened	Smoothened	Smoothened	Smoothened	Smoothened			
	9834	9834	10457	10457	10457	11968		11968		11968	,	11968			14198	14198	14198	14198	1464	14641	1464	14641	16041	1604]	16041	16041	16599	16599	16599	16599	16599
	555	226	557	228	529	200		561		295		563	. •		564	265	266	267	268	269	220	571	572	573	1574	1575	1576	1577	1578	. 629	280

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Homo sapiens	Homo sapiens	Homo saplens		Homo saplens		Homo sapiens		supple solution	Homo sapiens		Homo sapiens		Homo sapiens		Homo sapiens	•	Homo saplens		Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	_	Homo sapiens	Homo soplens		Homo sapiens	•	Homo sapiens		Homo sapiens	acolano omon	SI IBIODS OLLION	Homo saplens
EAEISPELØKRIGRKK	ANVTIGLPTKOPIPDC	SNASDSGSTQLPAPLR		CVLGYTELPADRAYVV		LNTVRKNAVRVHNØSD		NV PEIKIIKIKI KIRIN VIO	DSLDLRQLTRAGLRRL		EDADAENSSFYYYDYLDE		DKYLEIVHAQPYHRLRTR		CVLVRLRPAGGGRALK		DLGERQSENYPNKEDVGNK		EKLTKRLKRHPEETGGFQEA	KKEEKKEWRKTLEPWK	DPLHRTIETFAKEEPKEDID	YEIEYVCRGEREVVGPKVRK	SLWETV@KWREYRR@C		LOKDNSSLPWRDLSEC	CIVVSKIKANIMOKID		RWRLEHLHIGRDSSMKPLKC	•	CQVDETEEPDVHLPQP		REGLEAAGAAGASASYSS	los and violation	PLOCININGNICOLIN	ESKSSIKRVLAITTVLS
. 2670	2671	1227		1228		1249	טבטו	7/7	1273		363		364		365		3998		188	189	261	161	1205		1206	1208		1209		1520		1521	1500	1022	1523
NP_005622.1	NP_005622.1	043898		O43898		043898	073670	042090	043898		LR13		LR13		LR13	٠.	LR13		095375	095375	O95375	095375	AAA17021.1		AAA17021.1	AAA17021 1		AAA17021.1		NP_057456.1		NP_057456.1	ND 057464 1	100/400 L	NP_057456.1
Smoothened	Smoothened	G Protein-Coupled	Receptor GPR45	G Protein-Coupled	Receptor GPR45	G Protein-Coupled	Receptor GPR45	Receptor GPR45	G Protein-Coupled	Receptor GPR45	G Protein-Coupled	Receptor D6	G Protein-Coupled	Receptor D6	G Protein-Coupled	Receptor D6	G Protein-Coupled	Receptor D6	Gaba(b) Receptor 1	Gaba(b) Receptor 1	Gaba(b) Receptor 1	Gaba(b) Receptor 1	Glucagon-Like Peptide 1	Receptor	Glucagon-Like Peptide 1	Receptor Glucadon-Like Peptide 1	Receptor	Glucagon-Like Peptide 1	Receptor	G Protein-Coupled	Receptor LOC51210	G Protein-Coupled		Griolell Foodpled	G Protein-Coupled
16599	16599	17250		17250		17250	טאטענ	7 7	17250		17345		17345		17345		17345		17535	17535	17535	17535	17666		17666	17666		17666	٠,	18471		18471	18471	- 71	18471
1581	1582	1583		1584		1585	1504	2	1587		1588		1589		1590	•	1591		1592	1593	1594	1595	1596		1597	1598		1599		1600		1601	1400	3	1603

		02,00							40	7/448	3 -						.1,05	01,20		
	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homos caplens	
	QGTLEILYPDAHLSAED	PKTPLKERISLPSRRS	SVVQLRRQRPDFEWNEGLC	PAVGWHDTSERFYTHGC	AVQVGRQADRRAFTVPT	EHEPAGEEALRQKRAVATK	ALROKRAVATKSPTAE	CEKEVLSSNVSWRYEEQQLE	RLANNTGGWDSSGCYVEEGD	CKQEKSSLFQISKSIG	CIAFQRREGGVPGTRPGSPG	APGTRASRRCDRAGRWE	CPAERVANNRGDFRWPR	QNPPPEPPADQQLRFRC	VPLGGGAPGTRASRRC	PAARVHRPSRCRYRD	TLARPDATQSQRRRKTVRL	RSKLVAASVPARDRVRG	ACSERSAVITOATRPD	
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	1524	1525	2030	2032	2047	1513	1514	1515	1518	1519	2164	2166	2167	1712	2175	425	426	427	428	ì
-	NP_057456.1	NP_057456.1	ENSP00000164265	ENSP00000164265	ENSP00000164265	G9UIZ3	Q9UIZ3	ezin60	Q9UIZ3	Q9UIZ3	BAA96055.1	BAA96055.1	BAA96055.1	BAA96055.1	BAA96055.1	1729	U229	6221	LR29	
		•	•						•								663	<u>%</u>	<u>8</u>	663
Deceptor 10051010	G Protein-Coupled Recentor I OC51210	G Protein-Coupled Receptor I OC51210	G Protein-Coupled Receptor 1,819072	G Protein-Coupled Receptor 1s 19072	G Protein-Coupled	G Protein-Coupled	G Profein-Coupled	G Protein-Coupled	G Protein-Coupled	Receptor KIAAU/58 G Protein-Coupled	Receptor KIAA0758 G Protein-Coupled	Receptor Ls21632 G Protein-Coupled	Receptor GPR92/GPR9 G Protein-Coupled	Receptor GPR92/GPR9 G Protein-Coupled	Receptor GPR92/GPR9 G Protein-Coupled	Receptor GPR92/GPR9				
	18471	18471	19072	19072	19072	19501	19501	19501	19501	19501	21632	21632	21632	21632	21632	22315	22315	22315	22315	} ·
	1604	1605	9091	1607	1608	1609	1610	1611	1612	1613	1614	1615	1616	1617	1618	1619	1620	1621	1622	

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Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo conjens		Homo saplens	Homo saplens	! 	Homo saplens	Homo sapiens		Homo saplens	Homo socions		Homo saplens		Homo sapiens	•	Homo sapiens		Homo saplens	Homos omoH		Homo sapiens		Homo saplens		Homo sapiens		Homo sapiens		Homo sapiens
CSGKSTESSIGSGKTSGSR	ENTISTOTEDPDAKC	SSASLNREGLLNNARD	DRYIKINRSIQQRKAIT	CEHVEDKHNAKGEAIEN		RISKRRSKFPNSGKYA	CQLLFRRFQGEPSRSESTSE		RLQEIILTFEKINKTR	KGKSRAAENASLGPTN		LLFGTIMDHKIRDALR	IdWII/V\UGSXSSEJISdd		KLPNNELHGGESHNSGN		SGNRSDGPGKNTTLHNEFD		ROFISOSSRKRKHNOSIR		SHLDRLLDESAQKILYYC	CRSESPRI EKKSNIPTESE		ESIRSLQSVRRSEVRIYYD		CRKELSNLTEEEGGEGGV	3	EEDAGRIGRKNSSTSTSSS		CFGDRYYREPFVQRQRTSR	INOSCOSOS OT COTSSH	TOSS GDI GEO CONTROLL
1138	1141	1497	1255	1257		1258	1259	•	2721	2722	0	2/23	2724	i	1579		1580		1581	•.	1582	1584		1585		331	•	332		333	758	1,00
094867	094867	094867	095853	095853		095853	095853	1	CAC27252.1	CAC27252.1		CAC2/252.1	CAC27252.1		NP_076404.1		NP_076404.1	-	NP_076404.1		NP_076404.1	NP 076404 1		NP_076404.1		075963		075963		075963	075043	360
Latrophilin-3	Latrophilin-3	Latrophilin-3	G Protein-Coupled	G Protein-Coupled	Receptor GPR34	G Protein-Coupled	Receptor GPR34 G Protein-Coupled	Receptor GPR34	G Protein-Coupled Receptor L30698	G Protein-Coupled	Receptor Ls30698	G Protein-Coupled	Receptor Ls30698 G Protein-Coupled	Receptor Ls30698	G Protein-Coupled	Receptor GPR87/GPR95	G Protein-Coupled	Receptor GPR87/GPR95		Receptor GPR87/GPR95		Receptor GPR8//GPR45 G Protein-Coupled	Receptor GPR87/GPR95		Receptor GPR87/GPR95	G Protein-Coupled	Receptor RE2	G Protein-Coupled	Receptor RE2	G Protein-Coupled	Receptor REZ G Protein-Counted	
22925	22925	22925	25359	25359		25359	25359		30698	30698	00,00	3005	30698		30875		30875		30875		30875	30875		30875		31568		31568		31568	31568	3
1623	1625	1626	1627	1628		1629	1630		<u>8</u>	1632	007	3	1634		1635	٠.	1636		1637		1638	1639		1640		<u>3</u>		1642	:	1643	1644	<u>F</u>

	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	
	COKLOKIDLRHNEIYEIKVD	NKGDNSSMDDLHKKDA.	QDERDLEDFLLDFEED	ERGFSVKYSAKFETKA	RSKHPSLMSINSDDVEKQSC	DAQKESTGVTLRQRR	CKKINQLISETEAVVTN	ADDQTLLEQMIMDQDDG	KYNGSISLRRPRLASQ	KRYFAKFEEKFFQTC	DGDRQKAMKRLRVPPL	RVRSGRVRSYSTRDFQDC	CNNSVPGKEHPFDITVMIRE	APSKPGLPKPQATVPRKVD	AASKPKSTPAVIQGPSGKD	KRSELNKTLQTLSETYFIMC	GNASTERNGVSFSVQNGDVC	CRIKKKKQLGAQRKTSIQD	DFTGKQHMFNEKEDSC	
	1232	1233	1234	1235	1236	2597	2600	2610	2672	2673	2674	2103	2105	2106	-2135	1261	1262	1263	1264	
. *	075473	075473	075473	075473	075473	NP_004727.1	NP_004727.1	NP_004727.1	NP_004727.1	NP_004727.1	NP_004727.1	CAC28410.1	CAC28410.1	CAC28410.1	CAC28410.1	000406	000406	000406	000406	
	Receptor RE2 G Protein-Coupled Receptor GPD40	pe -	p _e	D	G Protein-Coupled	Xenotropic and Polytropic	Kenotropic and Polytropic Defroving Becomes (XPR)	Xenotropic and Polytropic Detroving Receptor (XPR)		Retrovirus Receptor (XPR1) Xenotropic and Polytropic	Retrovirus Receptor (XPR1) Xenotropic and Polytropic	Retrovirus Receptor (XPR1) Lung Seven Transmembrane	receptor z (Lustikz) Lung Seven Transmembrane December 2 (Histor)	Lung Seven Transmembrane Receptor 2 (118792)	Lung Seven Transmembrane	G Protein-Coupled	Receptor Grigot G Protein-Coupled Becentor GPD64	G Protein-Coupled	G Protein-Coupled Receptor GPR64	
	36534	36534	36534	36534	36534	37498	37498	37498	37498	37498	37498	40881	40881	40881	40881	42697	42697	42697	42697	
	545	946	647	648	646	920	. 159	652	653	23	655	939	657	658	629	98	[99	662	663	

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Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homos omoh		Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens
PNVNPASAGNQTQKTQD RVKSPPFAGTQI PKIIFS	KDGYMVVNVSSISINEPED	RSTVDSKAMGEKSFSVHNNG	COPLRARSLLTPRRTR	GQKHELETADGEPEPASRVC	KKTFIQGGQVSLVRHKD	CGEHHPMKRLPPKPQSP	STSTPGSSTPSRLELLSEE	METSSPRPPRPSSNPG	CSQVPSTSTPGSSTPSR	DPNGNESSATYFILIG	RHATVLTLPRVTKIGV	ILKTVLGLTREAQAKA	HRFSKRRDSPLPVILAN	KEIRORII RI FHVATHASF		GEDIEISDTESFSNDPC	SSKQIKTISGKTPQQYE	AATQNRRFQFTQNQKKE	CKDPIEDINSPEHIQRR •	CVLSRKIQEEYYRLFKNVP	CIAANINKTLTKIRSIKEP	KLSVNHRRTHLTKLMHTVE	EKITFLSHRKVTDRYRSLC	SSSLLGYKNNTISAKD	CSSYELQQQSMKRSNRRK
		•									٠			•										,	
2072	2074	2076	1265	1266	1267	1269	2294	2301	2302	1850	1851	1852	1853	1854	}	1416	1417	1419	1420	2113	2114	2115	2116	2117	1421
AAK57695 AAK57695	AAK57695	AAK57695	095665	095665	095665	095665	095665	095665	095665	1R76	LR76	LR76	LR76	LR76		075899	075899	075899	075899	NP_071442:1	NP_071442.1	NP_071442.1	NP_071442.1	NP_071442.1	P20309
KIAA1624 Protein KIAA1624 Protein	KIAA 1624 Protein	KIAA 1624 Protein	Neurotensin Receptor type 2	Neurotensin Receptor type	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	Receptor L353440 G Protein-Coupled	Receptor LS53440 G Protein-Coupled	Receptor LS53440	Gaba(b) Receptor 2	Gaba(b) Receptor 2	Gaba(b) Receptor 2	Gaba(b) Receptor 2	ETL protein	ETL protein	ETL protein	ETL protein	ETL protein	Muscarlnic acetylcholine					
45937 45937	45937	45937	50847	50847	50847	50847	50847	50847	50847	53440	53440	53440	53440	53440		54053	54053	54053	54053	55728	55728	55728	55728	55728	56923
1664 1665			1668	1669	1670	1671	1672	1673	1674	1675	1676	1677	1678	1679		089	1681		•			1686	1687	1688	1689

WO	02/00	91087						411	1/448	•					PCI		01/:	2016	,,	
Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens		Homo sapiens	Homo sapiens		Homo saplens		Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens
KPSSEQMDQDHSSSDSWNNN	DLERKADKLQAQKSVD	KEATLAKRFALKTRSQ	PPTCRPRRMSVCYRPPGNE	CLAVTRPFLAPRLRSPALAR	RGARWGSGRHGARVGR	TAGDLLPRAGPRFLTR	EGSGEARGGGRSREGTME	RTTPQLKVVGQGRGNGD	RSAPTALSRRLRARTHLPGC	VRGSHGEPDASLMPRSC		RKEDSVLMEA1SGGP1SFR	DQNKADIGGMLPGLTVRSV		PAGWPDQSLAESDSEDPSG		ETNHSLGKDDLRPSSP	SLVHELSGRRWQLGRRLC	LLYGWGEIYSEGSEEC FDV/GSDKTNSVSDISE	RHATVIFQPEGDTWREQK
1422	1423	1424	2097	2098	2099	2100	2101	2102	1909	0161	Ş	<u> </u>	1912		1913		2118	2119	2120	2122
P20309	P20309	P20309	NP_062813.1	NP_062813.1	NP_062813.1	NP_062813.1	NP_062813.1	NP_062813.1	NP_055061.1	NP_055061.1		NP_USSUOI.I	NP_055061.1		NP_055061.1		NP_076917.1	NP_076917.1	NP_0/091/. NP_0/091/.	NP_076917.1
Receptor M3 Muscarinic acetylcholine	Muscarinic acetylcholine Recentor M3	Muscarinic acetylcholine Pecenter M3	Leukofriene B4 Receptor Ri 1727	Leukotriene B4 Receptor	Cadherin EGF LAG Seven- Pass G-Type Percentar 1	(CELSR)/Flamingo) Cadherin EGF LAG Seven-	Pass G-Type Receptor 1 (CELSR1/Flamingo)	Cadherin Est LAG seven- Pass G-Type Receptor 1	(CELSR1/Flamingo) Cadherin EGF LAG Seven-	Pass G-Type Receptor 1 (CELSR1/Flamingo)	Cadherin EGF LAG Seven-	CELSR1/Flamingo)	5-HT5A Receptor	5-HT5A Receptor	5-HT5A Receptor	5-HT5A Receptor				
56923	56923	56923	57180	57180	57180	57180	57,180	57180	73584	73584	7000	73004	73584	2 1 2	73584	• .	74514	74514	74514	74514
1690	1691	1692	1693	1694	1695	1696	1697	1698	1699	1700		5	1702		1703		1704	1705	1707	1708

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PCT/US01/50107

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Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens		Homo sapiens	Homo sapiens	Homo sapiens		nomo sapiens	Homo sapiens		Homo saplens		sualdps office	Homo sapiens	Homo sapiens		Homo sapiens	Homo sapiens		Homo sapiens		Homo saplens	Homo sapiens	Homo sapiens	Homo saplens
GITRPESRPAVASQRR CHVVHGGEAAQQRPDDSEVE	RNPPAMSPAGQLSRTIE	RRLQPRLSTRPRRVSLC	RYLSVVSPLSTLRVPTLRC	SSILDTIFHKVLSSGCDYSE	VEILRTLFRSRSKRRHRTVK		GILFRIGIIIRSCEARGGLE	RLQAPSPASIPHSPGAFAYE	RIEPYYSIYNSSPSQEE		IIVIIAAILIKKINAAVIKKO	RNGNYNKLQHVQTRGYTKS		SRLQLVSAINLSTAKD	CKOKTB! DAMCKGN! EVND	CAGNINCIA DIVIDA GINE VINIA	NSAYMLSPKPQKKFVDQAC	CKVQDSNRRKMLPTQF		HAVSLIKLVIKGIKKPLS	NVNVFSELSAPRRNED		TKQRNPMDYPVEDAFC		CKPQLVKKSYGVENRA	RRAVPGHQAHGANLRH	KEDKLELTPTTSLSTRVNRC	KETLFMAGDTAPSEATSGEA
1277	1279	1280	155	. 156	157	ć.	80.	159	1589	1500	060.	1591		1592	1503	2	1594	1218		. 6171	1220		1221		1222	1286	1287	1288
P21731 P21731	P21731	P21731	AAA62837.1	AAA62837.1	AAA62837.1	1 7 6 0 0 7 4 4 4	AAA02837.1	AAA62837.1	NP_006785.1	NP OOK785 1	-000,000	NP_006785.1		NP_006785.1	NP 006785 1		NP_006785.1	AAC98506.1		AAC	AAC98506.1		AAC98506.1		AAC98506.1	AAB05897.1	AAB05897.1	AAB05897.1
Thromboxane A2 Receptor Thromboxane A2 Receptor	Thromboxane A2 Receptor	Thromboxane A2 Receptor	Chemokine (C motif) XC Receptor 1 (CCXCR1)	Chemokine (C motif) XC	Chemokine (C motif) XC	Receptor 1 (CCXCR1)	Receptor 1 (CCXCR1)	Chemokine (C motif) XC	G Protein-Coupled	Receptor GPIX/5 G Protein-Counled	Receptor GPR75	G Protein-Coupled	Receptor GPR75	G Protein-Coupled	G Protein-Coupled	Receptor GPR75	G Protein-Coupled Recentor GPD75	G Protein-Coupled	Receptor RAIG1	Gerentor RAIG1	G Protein-Coupled	Receptor RAIG1	G Protein-Coupled	Receptor RAIG1	G Protein-Coupled Receptor RAIG1	Tachykinin Receptor 2	Tachykinin Receptor 2	Tachykinin Receptor 2
81765 81765	81765	81765	98519	98519	98519	08510	\$100 \$	98519	130108	130108	2	130108		130108	130108	3	130108	133117	71001) [20]	133117	-	133117	1,1001	13311/	152198		152198
1709	1711	1712	1713	1714	1715	7171	2 .	7171	1718	1719	:	1720		1721	1722	 	1723	1724	3051	24	1726		1727	פטרי	9 7/1	1729	1730	1731

Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens		Homo sapiens		Homo sapiens		Homo sapiens		Homo sapiens	Homos cmoH		Homo sapiens	· •	Homo sapiens		Homo sapiens		Homo sapiens	Homo sopiens	200
CVVAWPEDSGGKTLLL RORKSVNAI NSPI HOF	KFQDTHNNAHYYVFFEEQED	CHVKIYITVRNPQYNPGDK	CKRQAQAYRGQRVPPKNSTD	SRSRFIRNTNESGEEVTT	COKEDSVYVCGPYFPRGWNN	SGEEVTTFFDYDYGAPCHKF	DFDDLNFTGMPPADEDYSPC	CWGLSMNLSLPFFLFRQAYH	RHRVTSYTSSSVNVSSN	CMLETETLNKYVVIIAYALV	EEPTNISTGRNASVGNAHRQ	RIRNPFTVYITHLSIAD	YVMCIDREESHSRNDCRAV	SSTILVVKIRKNTWASHSSK	TRAFKDEMQPRRQKDNC	ERYLGVAFPVQYKLSRRPL		QYLNTTEQVRSGNEITC		EGINEDRGVGQGEGMPSSD		RGLQVLRNQGSSLLGRRGKD		KACLEEAQLENETIGCS	KDI AI EDSGESDOCSE		LQKLRPPDIRKSDSSP		NPKYRHPSGGSNGATC		KVFSNFYSKAGNISKNC		CGYSDPEDESKITFYI	KRKWRSRCPTPSASPD	
1290 1445	1446	1449	1450	1896	1898	1899	908	807	808	1490	1527	1528	1529	1530	1531	1578		1586		1588		1616		1292	1206	2	1297		1298		1299		1301	1305	
AAB05897.1 P16473	P16473	P16473	P16473	NP_000639.1	NP_000639.1	NP_000639.1	P25024	P25024	P25024	P25024	NP_002368.1	NP_002368.1	NP_002368.1	NP_002368.1	NP_002368.1	NP_005297.1		NP_005297.1		NP_005297.1		NP_005297.1		P32241	D302A1		P32241		P32241		P41587		P41587	P41587	
Tachykinin Receptor 2 Thyrotropin Receptor	Thyrotropin Receptor	Thyrotropin Receptor	Thyrotropin Receptor	C-C Chemokine Receptor 2	C-C Chemokine Receptor 2	C-C Chemokine Receptor 2	Interleukin-8 Receptor A	Interleukin-8 Receptor A	Interleukin-8 Receptor A	Interleukin-8 Receptor A	Mas Proto-Oncogene	Mas Proto-Oncogene	Mas Proto-Oncogene	Mas Proto-Oncogene	Mas Proto-Oncogene	G Protein-Coupled	Receptor GPR43	G Protein-Coupled	Receptor GPR43	G Protein-Coupled	Receptor GPR43	G Profein-Coupled	Receptor GPR43	Vasoactive Intestinal	Polypeptide Receptor 1 Vasacetive Intestinal	Polypeptide Receptor 1	Vasoactive Intestinal	Polypeptide Receptor 1	Vasoactive Intestinal	Polypeptide Receptor 1	Vasoactive Intestinal	Polypeptide Receptor 2	Vasoactive intestinal	Polypeptide Receptor 2 Vasoactive Intestinal	
152198	152201	152201	152201	152245	152245	152245			152299	152299		158822	158822	158822	158822	159152		159152		159152		159152		159973	150073		159973		159973		160040		160040	160040) }
1732 1733	1734	1735	1736	1737	1738	1739	1740	1741	1742	1743	1744	1745	1746	1747	1748	1749		1750		1751		1752		1753	1754		1755		1756		1757	. (1758	1759	<u>}</u>

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	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens		Homo saplens		nomo sapiens	Homo sapiens	-	Homo sapiens	Homo sapiens	aclass omon	s jeiche oi jou	Homo sapiens	<u>.</u>	Homo saplens		Homo sapiens	Homo saplens		Homo saplens		Homo sapiens		nomo sapiens	Homo sapiens	Homos capings	
G. 17 (12 (14 G2133))	CGSSFSKINGSEGALAFFIK	REPPWPALPPCDERRCS	SPPSGPETAEAAALFSREC	SSRRPLRGPAASGRERGHRQ	RKSRPRGFHRSRDTAG	NPLVTGYLGRGPGLKTVC	•	GRYLGAAFPLGYQAFRRPC		CLEAWORASAGIRAS	CLRALARSGLTHRRKLR		NASNVASFLYPNLGGSWRK	TVSLPLKAVEALASGA	O/(dsOly/divio) istnsho		CSEAFPSRALERAFALY		ERAGAVRAKVSRLVAAVV		KKPGPSDPAAPHAELHKLGS	GAPANASGCPGCGANASD		DLFNHTLSECHVELSQST		NVLTACRURGPGGPKSRRHC		KDGIRAGICASSSCSIG	KGDSQPAAAPHPEPSLS	A IIVHIN IVI ODA DODOGOVIVI IN	
7001	000	132	134	135	136	1595		1596	1607	/ <u>40</u>	1598		1599	1617	1418	2	1926		1927	0001	1928	1929		390		391	Ç	760	484	7201	
041607	75.14	AAC26081.1	AAC26081.1	AAC26081.1	AAC26081.1	NP_005294.1		Receptor NP_005294.1	Docomtor NB 006004.1	INF_000294.1	Receptor NP_005294.1		Receptor NP_005294,1	Receptor NP_005294.1	Pacentor NP 005204 1	1.1.2000	BAB55446		BAB55446	0 0 0 0 0 0 0 0	pAp30440	BAB55446		015218		015218	010310	013210	015218	1885	
Polypeptide Receptor 2	Polypeptide Receptor 2	Motilin Receptor (GPR38)	Motilin Receptor (GPR38)	Motilin Receptor (GPR38)	Motilin Receptor (GPR38)	G Protein-coupled Receptor		G Protein-coupled Receptor			peldnoo-ule		G Protein-coupled Receptor GPR40	ein-coupled	in-compled		G Protein-Coupled	Receptor GPR54	G Protein-Coupled	Receptor Grissa	G Florell F-Coupled Receptor GPR54	G Protein-Coupled	Receptor GPR54	Adrenomedullin Receptor	(ADMR)	Adrenomedullin Receptor		(ADMR)	Adrenomedullin Receptor	(ADINIT) G Protein-Coupled	Receptor RTA
1,60040	9	160055	160055	160055	160055	160059	1	160059	140050	6000	160059		160059	160059	160059		160189		160189	001071	60,001	160189		160202		160202	140202	100202	160202	160204	
1760	3	1761	1762	1763	1764	1765		1766	1747		1768		1769	1770	1771	•	1772		1773	1774	1/1	1775		1776	. !	1777	1778	2	1779	1780)

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Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
CPGLSEAPELYRRGFLTIEQ	RDGAELGEAGGSTPNTVT	LAGRDKSQRLWEPLRV	RTIRKWNGCTHCYLAFNSD	RAKLLREGWVHANRPKR	RRVMLKEIYHPRMLLI	SALARAFGEEFLSSC	RSCSRKMNSSGCLSEE	PGPDRDATCNSRQAALAVSK	SSHAAVSLRLQHRGRRRPGR	DDSELGGAGSSRRRRTSSTA	DGPPEPGAEQHLELEPGPRR	CPILEQMSRLQSHSNTSIRY	RYIDHAAVLLHGLASLLGLV	CRMRQTVVTTWVLHLALSDL	SASLPFFYFLAVGHSWE	CLVLWALAVLNTVPYFVFRD	CYYNVLLINPGPDRDAT	CNSRQAALAVSKFLLAFLVP	RGLPFVTSLAFFNSVANPVL
1983	1985	2173	1678	1679	1680	1682	1683	151	152	153	151	2220	2221	2222	2223	2224	2225	2226	2228
LR85	1.785	1885	NP_001497.1	NP_001497.1	NP_001497.1	NP_001497.1	NP_001497.1	AAD21055.1	AAD21055.1	AAD21055.1	AAD21055.1	NP_004769.1	NP_004769.1	NP_004769.1	NP_004769.1	NP_004769.1	NP_004769.1	NP_004769.1	NP_004769.1
G Protein-Coupled	G Protein-Coupled Receptor RTA	G Protein-Coupled Receptor RTA	G Protein-Coupled	G Protein-Coupled	Receptor GPR32 G Protein-Coupled	Receptor GPR32 G Protein-Coupled	G Protein-Coupled Recentor GP032	G Protein-Coupled	G Protein-Coupled	Receptor GPR44 (CRIHZ) G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	Receptor GPR44 (CRIHZ) G Protein-Coupled	Receptor GPR44 (CRTH2) G Protein-Coupled	Receptor GPR44 (CRIHZ) G Protein-Coupled	Receptor GPR44 (CR1HZ) G Protein-Coupled December (CR1HZ)	G Protein-Coupled	Receptor GPR44 (CRIHZ) G Protein-Coupled
160204	160204	160204	160206	160206	160206	160206	160206	160210	160210	160210	160210	160210	160210	160210	160210	160210	160210	160210	160210
1781	1782	1783	1784	1785	1786	1787	1788	1789	1790	1791	1792	1793	1794	1795	1796	1797	1798	1799	1800

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	Homo sapiens	Homo sapiens		nomo sapiens	Homo sapiens		sueidos outou	Homo saplens	Homos carolens		Homo sapiens		Homo sapiens	Homo sapiens		Homo sapiens	Homo saplens		Homo saplens	Homo sapiens		Homo sapiens	Homo saniens		Homo sapiens		ivius musculus	Homo sapiens	
	CSRPEEPRGPARLLGWLLGS	CAASPQTGPLNRALSS	KEINIDBBADEDGUEVASODE	BY COUNTY OF THE WAY O	CVKDQEAQEPKPRKRANS			HSCPLGFGHYSVVDVCIFE	GKVEKYMCEHNMSDDTWSAK		RSIHILLGRRDHTQDWVQQK		CKAKGSISFFLGLSIM	KEFRMNIRAHRPSRVQLVLQ		AGIRPIDVGGAEAIIKAAIK	KEFQEASALAVAPRAKAHK		GGFCFRSTRHNFNSMR	ETIRRALYITSKLSDANC		FFF VILLIGG GUDEDAP CALES	RGARRLLVLEEFKTEKRLC		NASEPGGSGGEAAALGLK		というとこうとこうとうとうとう	RPAGPGRGARRLLVLE	
	2229	2230	444	Į.	445	776	2	622	161		162	149	<u>3</u>	164	c	7	· •	-	123	125		coo	338		496	אוצ	2	1291	,
		_		.•					_					_	_	_	_											• .	
	NP_004769.1	NP_004769.1	Q9Y2T5) }	Q9Y2T5	COVOTS)	Q9Y2T5	AAD22410.1		AAD22410.1	4 A D 20010 1	VVV24410.	AAD22410.1	A A CESCOS 1	יייייייייייייייייייייייייייייייייייייי	AAC52028.1	• .	AAC52028.1	AAC52028.	2	2	927		126	0.54807		- 921	
,,,,	Receptor GPR44 (CRTH2) G Protein-Coupled	Receptor GPR44 (CRIHZ) G Protein-Coupled	Receptor GPR44 (CRTH2) G Protein-Coupled	Receptor GPR52	G Protein-Coupled	Receptor GPR52 G Protein-Coupled	Receptor GPR52	G Protein-Coupled	Receptor Grigoz G Protein-Coupled	Receptor GPR55	G Protein-Coupled	Receptor GPR00	Receptor GPR55	G Protein-Coupled	Receptor GPR55 G Protein-Coupled	Receptor GPR35	G Protein-Coupled	Receptor GPR35	G Protein-Coupled Receptor GPR35	G Protein-Coupled	Keceptor GPR33	Receptor GPR27	G Protein-Coupled	Receptor GPR27	G Protein-Coupled	G Protein-Coupled	Receptor GPR27	G Protein-Coupled	ונפכפים סבונדי
	160210	160210	160212		160212	160212		160212	160217		160217	160217	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	160217	01/210		160219		617091	160219	1,60001	7700	160221		160221	160221		160221	
	1801	1802	1803		1804	1805	1	908	1807	6	808	1800	}	1810	1811	: ·	1812		8 8 8 8	1814	1815	2	1816		/ 18 18	1818		1819	

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Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homos canon		Homo saplens	Homo saplens	Homo saplens	5	Homo sapiens		SI DICTOR OLLOW	Homo saplens		Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens		Homo sapiens	
CQRPPKPQEDGQPSPV	CNMIGDVTTEQYFALRRK	EGRADEQSAEAALAVP	GNFVGRRRYGAESQNPTVK	RIFRSIKOSMGLSAAQKAK	CDREVAVAVALESPOR		ATDHSRQEVSRIHKGWKE	KTDVTRLTHSRDTEELQS	FTOFOOSPSKRGTFDFFAK		SPNPDKDGGTPDSGQELR		C&LV I VVI V I CORNOL	AANGSDNKLKTEVSS		PRDSFRGSRSLSFRMRE	ERFATMVRPVAESGATKTSR	RLVQASGQKAPRPAAR	RAVEAHSGASTTDSSLRPRD	IFRLVQASGQKAPRPAAR	DSSLRPRDSFRGSRSLSFRM	RSLSFRMREPLSSISSVR	GPEDGGLGALRGLSVAASC	ANIGSLCVSFLQPKKE •	ETIFNAVMLWEDETVVE		CNRKVYQAVRHNKATENKE	
1606	1607	1610	1611	1600	1,601	3	1604	1605	403		404	30%	}	406		۵,	71	72	73	1914	1915	1916	1917	1625	 1626		1627	
NP_057624.1	NP_057624.1	NP_057624.1	NP_057624.1	NP_037477.1	NP 037477 1		NP_037477.1	NP_037477.1	060883		O60883	0,60,883		060883							CAA04118.1	CAA04118.1	CAA04118.1	NP_003599.1	 NP_003599.1	1	NP_003599.1	
G Protein-Coupled Receptor GPR72	G Protein-Coupled Receptor GPR72	G Protein-Coupled Receptor GPR72	G Protein-Coupled	receptor GPIK/2 G Protein-Coupled	Receptor G2A G Protein-Coupled	Receptor G2A	G Protein-Coupled Receptor G2A	G Protein-Coupled	Receptor G2A Endothelin Type B Receptor-	Like Protein 2 (ETBR-LP-2)	Endothelin Type B Receptor-	uke riolem z (Elbik-Lr-z) Endothelin Ivne B Recentor-	Like Protein 2 (ETBR-LP-2)	Endothelin Type B Receptor-	Like Protein 2 (ETBR-LP-2)	Sphingolipid Receptor Edg6	Sphingolipid Receptor Edg6	Sphingolipid Receptor Edg6	Sphingolipid Receptor Edg6	Sphingolipid Receptor Edg6	Sphingolipid Receptor Edg6	Sphingolipid Receptor Edg6	Sphingolipid Receptor Edg6	T-Cell Death-Associated	T-Cell Death-Associated		T-Cell Death-Associated	Gelie o (GPIKOS)
160222	160222	160222	160222	160223	160223	•	160223	160223	160224		160224	160224		160224							160225	160225	160225	160228	160228		160228	•
1820	1821	1822	1823	1824	1825	,	1826	1827	1828	-, -!	1829	1830	} .	.1831		1832	1833	1834	1835	1836	1837	1838	1839	1840	1841		1842	

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Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens		Homo saplens		Homo sapiens	Homo sapiens	•	Homo sapiens	Homo saniens	Homo saplens	Homo saplens	Homo sapiens			Homo sapiens			Homo saplens			Homo sapiens	
CILEHAVNFEDHSNSGKR	CNTSQRQRKRILSVSTKD	CDAEKSNFTLCYDKYPLEK	CTVDWKSKDANDSSFV	CVEDLQTIQVIKILKYEK	CORPAKDLPAAGSEMQIRP	TSDESLSVDDSDKTIG	ERHVAIAKVKLYGSDKSC	RSRDLRREVLRPLQC	QEHYNYTKETLETQET	GRRRVGTPGHHLLPLR	MMRKKAKFSLRENPVEETKG		MMIEYSNFEKEYDDVTIKM		CEQTEEKKLKRHLALFRSE	KKRVGDGSVLRTIHGKEMSK		DRARRERFIMINEKWDTNSSE	RKNGEQWHVVSRKKGKIIK	RKSAEKPQQELVMEELKE	RQSAGDRRRLGLSRQTAK	DRFLKIIRPLRNIFLKKP			MILSNKEATPSSVKKC			VYDSYRKSKSKDRKNN			AKARATINGTINGKIDO	
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1628	1629	2303	2131	2132	2133	2134	1018	1019	1020	1021	1922	<u>:</u> 	1923		1924	1925		463	464	465	200	1619			1620			1622		1409	200	
NP_003599.1	NP_003599.1	NP_003599.1	NP_055137.1	NP_055137.1	NP_055137.1	NP_055137.1	095136	095136	095136	095136	ENSMPRT221753		ENSMPRT221753		ENSMPRT221753	ENSMPRT221753		Q9Y5X5	Q9Y5X5	Q9Y5X5	Q9Y5X5	NP_076403.1			NP_076403.1			NP_076403.1		ND 074409 1	1.00400.1	
T-Cell Death-Associated Gene 8 (GPR65)	T-Cell Death-Associated Gene 8 (GPR65)	T-Cell Death-Associated Gene 8 (GPR65)	Encephalopsin	Encephalopsin	Encephalopsin	Encephalopsin		Sphingolipid Receptor Edg5		Sphingolipid Receptor Edg5	0	Receptor GPR103	G Profein-Coupled	Receptor GPR103	G Protein-Coupled Receptor GPR103	G Protein-Coupled	Receptor GPR103	Neuropeptide FF 2 Receptor	G Protein-Coupled	Receptor	GPR86/GPR94/P2Y13	G Protein-Coupled	Receptor	GPR86/GPR94/P2Y13	G Protein-Coupled	Receptor	GPI386/GPI394/P2Y13 G Protein-Coupled	Pecentor Pecentor	GPR86/GPR94/P2Y13			
160228	160228	160228	160300	160300	160300	160300	160312	160312	160312	160312	160314		160314	.•	160314	160314		160317	160317	160317	160317	160324			160324			160324		1,40304	10001	
1843	1844	1845	1846	1847	1848	1849	1850	1851	1852	1853	1854		1855		1856	1857		1858	1859	1860	1861	1862			1863			1864		18.65	3	

Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens		Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	. Homo sapiens	Homo sapiens		Homo sapiens		
CMQGRKTTASSQENHSSQTD	CANDSDTLELPDSSRA	PLRARALRGRRLALGLC	LGRGTFRLARSDRVLC	RDKVRAGLFQRSPGDT	CELKRDLQLLSQFLKHPQK	TSVRFMGDMVSFEEDR	RQEEEQSEIMEYSVLLP	RILFORTKGRSGEAEKR		GSLLEETTRKWAQYKQAC	QTIENATDIWQDDSEC	CPKKLSEGDGAEKLRK	QQDHARWPRGSSLSEC	EPTSTHESEHQSGAWC	CEPREVRRYQWPATQQ	RSGDFPPGDGGPEPPR	CTAEDGATSRPLSSPPGRDS	RESAGKNYNKMHKRERTC	RDSPSYPDSSPEGPSEALP	QVGPCRSLGSRGRGSSGAC	•	CRDAGTELTGHLVPHHDGLR		
1624	1308	1309	1310	1311	1213	1214	1215	1216) - -	1312	1313	1315	1316	1121	1126	1129	1131	1706	1707	1938		1939	•	
				•	,	-			,															
NP_076403.1	076067	076067	076067	076067	Q9Y653	Q9Y653	Q9Y653	Q9Y653		095838	095838	095838	095838	094910	094910	094910	094910	094910	094910	NP_001399.1		NP_001399.1		
G Protein-Coupled	Receptor GPR86/GPR94/P2Y13 Protelnase-Activated	Receptor 4 Proteinase-Activated	Receptor 4 Proteinase-Activated Peceptor 1	Proteinase-Activated	G Protein-Coupled- Becoptor TM/7VN1 (CPD5.4	G Protein-Coupled-	Receptor IIVI/XINI/SPIKSO G Protein-Coupled-	Receptor TM7XN1/GPR56 G Protein-Coupled-	Receptor TM7XN1/GPR56	Glucagon-Like Peptide 2	Glucagon-Like Peptide 2	Receptor Glucagon-Like Peptide 2	Receptor Glucagon-Like Peptide 2 Popping	receptor Latrophilin-1	Latrophilin-1	Latrophilin-1	Latrophilin-1	Latrophilin-1	Latrophilin-1	Cadherin EGF LAG Seven- Pass G-Type Receptor 2	(CELSR2)	Cadherin EGF LAG Seven-	Pass G-Type Receptor 2	(CELSKZ)
160324	160329	160329	160329	160329	160330	160330	160330	160330		160387	160387	160387	160387	160388	160388	160388	160388	160388	160388	160390		160390	1	
1866	1867	1868	1869	1870	1871	1872	1873	1874		1875	1876	1877	1878	1879	1880	1881	1882	1883	1884	1885		1886		

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	WO 02/06	1087			420)/448		•			PCT/US0	1/5010)7
Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens Homo sapiens Homo sapiens Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens Homo sapiens Homo sapiens Homo sapiens Homo sapiens	Homo sapiens	Homo saplens
CKLAQAPGLRAGERSPEESL	RVSDTPEGVNSLDPSHGES	RSGKSQPSYIPFLLREES	CEALDSKGIKWPQTQR DILDAQLQELKPSEKD RTHSLLYQPQKKVKSE RDSPYPESSPDMEEDL CQEQKMLRTLDLSYNNIRD	CDSYANLNTEDNSLQD	KGTADAANVTSTLENEE	ERSLSAKDIMKNGKSNHLK	CNLEKEDLSENSQSSMIK	KRRVTKKSGSVSVSIS	CGTQSAHSDYADEEDS	DEEDSFVSDSSDQVQAC	ATILKLLRTEEAHGREGRR CRRVPRDTLDTRRESLFSAR PLSSKRWRRRRYAVAAC CRRMGPRSPSVIFMINL MMIPIKDIKEKSNVGC	CLVIRQLYRNKDNENYP	CSTRISLFKAKEATLL

1632

NP_060960.1

1631

NP_060960.

1633

NP_060960.1

1634

NP_060960.1

1635

NP_060960.1

1636

NP_060960.1

3 Protein-Coupled

1681

8

Receptor GPR48

3 Protein-Coupled

160411

190

eceptor GPR48

eceptor GPR48

1918 1919 1920

1280 1280

S160435 Receptor S160435 Receptor S160435 Receptor

160435

1902

60435

60435

60435

82 82

1637

NP_060960.1

1921

1225

1224

Platelet Activating Receptor 014626

lomolog (H963)

160889

1907

68809

160889

1908

lomolog (H963)

latelet Activating Receptor 014626

082 1780

S160435 Receptor

Platelet Activating Receptor 014626

1940

NP_001399.1

Cadherin EGF LAG Seven-Pass G-Type Receptor 2

160390

1887

1942

NP_001399.1

Cadherin EGF LAG Seven-

160390

1888

CELSR2)

Pass G-Type Receptor 2

CELSR2)

1943

NP_001399.

Sadherin EGF LAG Seven-

160390

1889

ass G-Type Receptor 2

095490 095490 095490 095490

atrophilin-2 atrophilin-2 atrophilin-2 atrophilin-2

160397 160397 160397 160397 160411

1890

89

1892 1893

CELSR2)

1136

. 096090 JN

3 Protein-Coupled

894

Receptor GPR48

3 Protein-Coupled

160411

1895

Receptor GPR48

3 Protein-Coupled

160411

1896

eceptor GPR48

3 Protein-Coupled

160411

1897

eceptor GPR48

3 Protein-Coupled

160411

1898

3 Protein-Coupled

16041

1899

eceptor GPR48

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,	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo saplens	***	Homo sapiens		Homo sapiens	Homo saplens		Homo sapiens	Homo sapiens	Homo caniens		Homo sapiens		Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Equine herpesvirus	2	
	ETFASPKETKAQKEKLRC	ESRAVGLPLGLSAGRRC	EDARGKRRSSLDGSESAK	RTVWEQCVAIMSEEDGD	CKVRFDANGATGPGSRD	RRLSHDETNIFSTPRE	GGPPEYLGQRHRLEDEED	REEITTFIDETPLPSP	RRPRPLGLSPRRLSLGSPE	RYGALELCVPAWEDARR	GAAAEARRRATGRAGR	ASRHFRARFRRLWPC	RARRALRRVRPASSGPP	ERYAAVLRPLDTVQRPKG	•	RAYRRSQRASFKRARRPGAR		KINYKUHUKGKVIKGPGGG	RARFQRCSGRSLSCSPQPTD		ARGHFDPEDLNLTDEALRLK	IGLRLRRERLLLMQEAKGRG	HOO IOULA A A DSDALLO IO		ALCLGACCHRLRPRHSS		CFFLLKPFRARDWKRRYD	PFPILRSTDLNNNKSC •	QLSRHGSSVTRSRLMSKE	LRQPPMAFQGISERQK	YYDDLDDVDYEESAPC		
	1226	1690	1691	1692	1693	1694	1695	1696	1697	202	203	204	205	37.1		372	c r c	5/5	374		394	395	306		397		859	860	862	863	1672		-
	014626	NP_062832.1	NP_062832.1	NP_062832.1	NP_062832.1	NP_062832.1	NP_062832.1	NP_062832.1	NP_062832.1	AAC35944.1	AAC35944.1	AAC35944.1	AAC35944.1	ાં ક	.•	ភោទ	i	חלוט	LR15		LR20	UZ20	0001		LR20		000398	000398	000398	000398	NP_042597.1	-	
Homolog (H963)	3 Receptor	Protein A	Protein A	Protein A	Protein A	Protein A	Protein A	Protein A	Protein A	Galanin Receptor GaIR3	Galanin Receptor GalR3	Galanin Receptor GalR3	Galanin Receptor GalR3	Urotensin-II Receptor	(GPR14)	Urotensin-II Receptor	(GPR14)	Urotensin-ii keceptor (GPR14)	Urotensin-II Receptor	(GPR14)	G Protein-Coupled Receptor GPR66	G Protein-Coupled	Receptor GPRes G Protein-Counted	Receptor GPR66	G Protein-Coupled	Receptor GPR66	Purinergic Receptor P2Y10	Purinergic Receptor P2Y10	Purinergic Receptor P2Y10	Purinergic Receptor P2Y10	G Protein-Coupled	Receptor Laid 1293 (Herpes	virus)
•	160889	161024			161024	161024	161024	161024	161024	161214	161214	161214	161214	161221		161221	.00.	17710	161221	• .	161249	161249	1612/0	†	161249		161251	161251	161251	161251	161293		
	1909	1910	161	1912	1913	1914	1915	1916	1917	1918	1919	1920	1921	1922	•	1923		1924	1925		1926	1927	1028	27.	1929		1930	1831	1932	1933	1934	:	

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Equine herpesvirus 2	Equine herpesvirus 2	Equine herpesvirus 2	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	•	Homo saplens	Homo sapiens
CDPYYPEMSTNVWRRAHVAK	CYYVIIRRLLRRPSKK	CKYIPFLSGDGEGKEGPT	RNLTSSPAPTASPSPAPS	PSWTPSPRPGPAHPFLQPP	RSSHQKRGTTRDVGSNVC	KSTSTTASFVSSSHMSVEE	TSSPFLMAKPQKDEKNNTKC	KKSMKKNLSSHKKAIG	QRTIHLHFLHNETKPC	RKHSLSSVTYVPRKKASLPE	RAVSYRAQGDTRRAVRK	GRRTRLRLDGAREAAGPE	GSFTQRFRLSRDRKVA	RYGVGEAAVGAEAGEATLG	SSRGTERPRSLKRGSKPSAS	KPSASSASLEKRMKMVS	RTTLFSFYFRDTPRANR	RPEMSRGLLAVRGAFV		CAVLSHRRAQPWALLLV	RVLVSDSLFVICALSL
1674	1675	1676	1820	1821	1822	1823	1317	1318	1319	1320	474	475	476	477	1477	1479	2052	2053	-	2059	2733
NP_042597.1	NP_042597.1	NP_042597.1	e NP_006670.1	e NP_006670.1	e NP_006670.1	e NP_006670.1	11 Q9Y271	11 Q9Y271	n Q9Y271	1 697271	Q9Y5N1	Q9Y5N1	Q9Y5N1	Q9Y5N1	Q9Y5N1	Q9Y5N1	NP_064540.1	NP_064540.1		NP_064540.1	NP_064540.1
G Protein-Coupled Receptor L3161293 (Herpes	virus) G Protein-Coupled Receptor Ls 1 6 1 293 (Herpes	G Protein-Coupled Receptor Ls 161293 (Herpes virus)	omedin K Receptor-Lik R)	Neuromedin K Receptor-Like NP_006670.1 (NK-4R)	edin K Receptor-Lik	Neuromedin K Receptor-Like (NK-4R)	yl Leukotriene CYSL or	Leukotriene CYSL	Leukotriene CYSL	Cysteinyl Leukotriene CYSLI	Receptor Histamine H3 Receptor	Histamine H3 Receptor	Histamine H3 Receptor	Histamine H3 Receptor	Histamine H3 Receptor	Histamine H3 Receptor	G Protein-Coupled Receptor OPEA	G Protein-Coupled	Receptor ORF4	G Protein-Coupled Receptor ORF4	G Protein-Coupled Receptor ORF4
161293	161293	161293	177147	177147	177147	177147	177168	177168	177168	177168	177191	177191	177191	177191	177191	177191	177387	177387	•	177387	177387
1935	1936	1937	1938	1939	1940	1941	1942	1943	1944	1945	1946	1947	1948	1949	1950	1951	1952	1953		1954	1955

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Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens		Homo saplens		Homo sapiens	Homo sopiens		Homo sapiens		sueidos ortion	Homo sapiens		Homo sapiens		Homo sapiens	Homo sanjens	
KRKTNVLSPHTSGSIS	CFSGENPERRPSRIPST	SYKDEDMYGTMKKMIC	VERHIVISIMRMRVHSN	CORMDIVIMKALALLAD	CSLRLPPERPRFAAFTAT	RGPLPPGICAHSAQGALRR	CRQAQARDLGAPWAVGLRSL	QQKLEDPFQKHLNSTEE	KKDKSLEADEGNANIQRPC	SQHDPQLPPAQRNIFLTEC	ILHPFRAKLOSTRRRALR	CKKRGTKTQNLRNQIRSK		EKPSSPSSGKGKTEKAE		PSVQDNDPIPWEHEDQETGE	KKPPTVSESGETPAGNSEG		LVMSEEFREGLKGVWK		GLTUN VTSTEST NSTEST	PDVEQFWHERDTVPSVQ		RHHEGVEMCLVDVPAVAEE		RVPQTPGPSTASGVPE	ETPRORSESUSSRSTMVTS	
1014	1015	1016	1017	443	528	533	534	420	422	423	487	415	•	418		419	486		1832	1000	200	1834		1835		1685	1686	
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AAF00530.1	AAF00530.1	AAF00530.1	AAF00530.1	LR37	LR37	LR37	LR37	LR28	LR28	LR28	LR28	LR27	•	LR27		LR27	LR27		1627		UK\$/	LR27		LR27		AAK12637.1	AAK12637.1	
Lysophosphatidic Acid	receptor Edg/ Lysophosphatidic Acid Pacantor Edg7	Lysophosphatidic Acid Receptor Edo?	Lysophosphatidic Acid	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled Receptor GPR78	G Protein-Coupled Receptor CPD78	Neuromedin U Receptor 2	G Protein-Coupled	Receptor Ls189884	G Protein-Coupled	Receptor Ls189884	G Protein-Coupled	G Protein-Coupled	Receptor Ls189884	G Protein-Coupled	Receptor Ls 189884	Receptor 1s 189884	G Protein-Coupled	Receptor Ls189884	G Protein-Coupled	Receptor Ls189884	G Protein-Coupled	Keceptor GPK6 I G Protein-Coupled				
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	Homo saplens	Homo sapiens	Homo saplens		Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens		Homo saplens			Homo sapiens		Homo sapiens	-		Homo sapiens		Homo saplens	<u>.</u>	Homo saplens	<u>.</u>	.Homo sapiens	Homo sapiens		Homo sapiens	Homo sapiens
	SSGAPQTTPHRTFGGGK	KPAPEEELRLPSREGSIEE	CPSESWVSRPLPSPKQE		ALERSLIMARRGPAPVSS	DGSFSGSERSSPQRDGLD	CGRDPSGSQQSASAAEASG	ASRKAEAIGKLKVQGEVS		SCLSYRVGTKPSASLR			KVDYYLLHETWRFGAAAC		HQSRALLGLTRGRQGPVSD			CIHTRPWTSNTVFLVSL		RGRQGPVSDESSYQPSR		IDRYLIKYPFREHLLQKKE		TDNGTICNDFASSGDPN •	FLKGRNRQVATALPLE		RNVRIASRLGSWKQYQC	GDHFRDMLMNQLRHNFKS
	1687	1688	1689	310	316	317	318	2266		2270			77/1		2272			2273	•	2274		2108		2109	2110		2111	2112
	AAK12637.1	AAK12637.1	AAK12637.1	[0]	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	ISI	l&I	ENSP00000071589		ENSP00000071589		1	EINSPUDDUDUD 1589		ENSP00000071589			ENSP00000071589		ENSP00000071589		AAK29080.1	1	AAK29080.1	AAK29080.1	•	AAK29080.1	AAK29080.1
Receptor GPR61	G Protein-Coupled Receptor GPR61	G Protein-Coupled	G Protein-Coupled	Receptor GPR61 Sobingolinid Receptor Edg8	Sphingolipid Receptor Edg8	Sphingolipid Receptor Edg8	otor Ed	G Protein-Coupled Recentor Is 180001	(HEOAD54)	G Protein-Coupled	Receptor Ls189901	(HECADS4)	G Moteur-Coupled Receptor Ls 189901	(HEOAD54)	G Protein-Coupled	Receptor Ls189901	(HEOAD54)	G Protein-Coupled	Receptor LS 189901 (HFOAD54)	G Protein-Coupled	Receptor Ls 189901	Purinergic Receptor P2U2	(GPR91)	Purinergic Receptor P2U2 (GPR91)	Purinergic Receptor P2U2	(GPR91)	Purinergic Receptor P2U2 (GPR91)	Purinergic Receptor P2U2 (GPR91)
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CVAFPLAVGNPDLQIPSR	NTLRHNALRIHSYPEGIC	QASKLGLMSLQRPFQMSID	DMIMPRSFKFLPGLPGH I KKIR QNLKDPVQIKIKHTRTQE	KNKSFGGWNTSGCVAHRD	CGRNGKRSNRTLREEVLR	TSKSKSSSTTYFKRNSHTD	DKSLSKLAHADGDQTS	LFPLLRTSDDTPGNRTKC QDKYPMAQDLGEKQKALK	SFPLDFLVKSNEIKSC	RRRLSRQDLHDSIQLHAK	KGEAKLDSRAKDVTLTIQE	DHKEQPIVTENAERQLVVKD	EGKEGDYIRIPERLIDVQD		
1721	1722	1723	1/24	1716	1717	1719	1720	407	409	410	1725	7271	1729		
AAK12639.2	AAK12639.2	AAK12639.2	AAK12039.2 Q9Y3K0	Q9Y3K0	@913K0	Q9Y3K0	Q9Y3K0	UR24 UR24	LR24	LR24	AAD55586.1	AAD55586.1	AAD55586.1		
G Protein-Coupled Receptor GPR63 (PSP24	beta) G Protein-Coupled Receptor GPR63 (PSP24 beta)	G Protein-Coupled Receptor GPR63 (PSP24 Deta)	G Protein-Coupled Receptor GPR63 (PSP24 beta) G Protein-Coupled	Receptor DJ28/914.2 G Protein-Coupled Receptor DJ287914.2 G Protein-Coupled	Receptor Dj287g14.2 G Protein-Coupled	Receptor Dj287g14.2 G Protein-Coupled Receptor Dj287g14.2	G Protein-Coupled Receptor Dj287g14.2	G Protein-Coupled Receptor JEG18 G Protein-Coupled	Receptor JEG 18 G Protein-Coupled	Receptor JEG 18 Receptor JEG 18	G Protein-Coupled Receptor VLGR1	G Protein-Coupled Receptor VLGR1 G Protein Coupled	Receptor VLGR1 G Protein-Coupled		
189920	189920	189920	189945	189945	189945	189945	189945	190026	190026	190026	190031	190031	190031		
1996	1997	1998	2000	2001	2003	2004	2005	2006	2008	2009	2010	2011	2013		

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	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens
	SEAYADGIEGYDILVACSSS	NNLRENGNNGVKKDKKAAK	DPFLNFSTPVVLFDALT	GKIFSSCFHNTILCMQKE	CPKFVNKILSSHQPLFS	KQHARVISHVPENTKGAVKK	ENTKGAVKKHLSKKKDRKA	CKFHTSFDMMLRLTSI	ENHDQDLDELQLEMEDSKP	NPHFRDDLRRLRPRAGDS	EDLHLDDEESSKRPLGLLAR	DSGPLAYAAAGELEKSSC	CAARROHALLYNVKRHSLE	DGSLKAKEGSTGTSESSV	CSIDLGEDGMEFGEDDIN	SEDDVEAVNIPESI.PPS •	MHKTIKKEIQDMLKKFFC	KEDSHPDLPGTEGGTEG	RQVKRAAQALDQYKLRQAS
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	324	326	379	380	327	328	329	330	439	044	442	. 129	1836	1837	1838	1839	1840	1841	343
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Receptor VLGR1	D ~	D.		þ	þ		pe	G Protein-Coupled Recentor GPD57	6	peld	peld	G Protein-Coupled	Gerapion Lerko G Protein-coupled Receptor CAC33098.	GPR101 GPR101	G Protein-coupled Receptor CAC33098.1	G Protein-coupled Receptor CAC33098.1	G Protein-coupled Receptor CAC33098.1	G Protein-coupled Receptor GPR101	nation-Related G Coupled Receptor
	190168	190168	190168	190168	190170	190170	190170	190170	190188	190188	190188	190188	190414	190414	190414	190414	190414	190414	190418
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	RTDEAMPGRFQELDSRLASG		DSSEVGDQINSKRAKQMAEK		KAQPIKGARRAPDSSSEFGK		RRKSNERLRGYSTGKT		RRGKSSYNYLLALAAAD	PALLACIA A A A A A A A A A A A A A A A A A A		CSIFFILNSIIVYKLR		GRLIYSLLSFISIPH	FFLFLWIHVDRE		MDPTISTLDTELTP	OSE/VS/ONCES COLUMNS A		RVLLKVEVPESGLRVSHRK		NINCH DE LEGISTATION OF THE PROPERTY OF THE PR	MEPNGTFSNNNSRNC		CTIENFKREFFPIVYLIF	GVLGNGLSIYVFLQPYK		ADYYLKGSINWIFGDLAC	FRLLHVTSIRSAWILC	
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	L78		LR8		L78		CAC33085.1		CAC33085.1	1 300000	(AC-30083.1	CAC33085.1		AAK91804.1	AAK91804.1		AAK91804.1	070		LR49		\	LR49		NP_065110.1	2 NP_065110.1	ני טרנשאט מוע	Z INP_000110.1	NP_065110.1	
EX33	٠: ٔ	Protein-Coupled Receptor EX33	nmation-Related G	EX33	Inflammation-Related G	Protein-Coupled Receptor	stein-Coupled	•••		Receptor LS 1904 19				Protein-Coupled	Receptor MrgX1 G Protein-Coupled	٠.	Protein-Coupled	Reception Cysteinyl Jenikotriene CVS 12	Receptor	Leukotriene CYSLT2	Receptor	۷.	Leukotriene CYSLT2	Receptor	Cystelnyl Leukotriene CYSLT2 NP_065110.1 Receptor	Leukotriene CYSLI	Receptor	Cystelriyi Leukoliiene Craliz Receptor	Cysteinyl Leukotriene CYSLT2 NP_065110.	
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	CGIIWILIMASSIMILIDSGS	CLELNLYKIAKLQTMNYIAL	VSHRKALTIIITLIIFFLC	CFLPYHTLRTVHLTTWKVGL	CKDRLHKALVITLALA	YFAGENFKDRLKSALRKG	HPGKAKTKCVFPVSVWLRKE	DSVSYEYGDYSDLSDRPVDC	RESQGQDESVDSKKSTSHD	PSAIYRRLHGEHFPARLGC	CHWALRESQGQDESVDSKKS	MGNDSVSVFVGDVSDI SDRPVDC	TERLIKIRWHISDNOVRPOAC	FADI CATCHIPPIDIFI DIFI		RTCHRQQQPAACRGFARVAR	EERPGSFTPTEPQTQLDSEG.	RSDPTAQPQLNPTAQPQSD		NAVIO I CICLE IN THE INTERIOR INTERIOR IN THE INTERIOR IN THE INTERIOR INTERIOR IN THE INTERIOR INTERI	KKKRMAMARRTMFQKGE	
	2257	2258	2260	2261	2262	2263	2264	429	430	431	432	2818	2585	434	t P	435	436	437	1730	2	1731	
	NP_065110.1	CYSLT2 NP_065110.1	CYSLT2 NP_065110.1	NP_065110.1	CYSLT2 NP_065110.1	CYSLT2 NP_065110.1	CYSLT2 NP_065110.1	LR31	LR31	เหลา	LR31	NP 060955.1	ENSP00000080322	1833		LR33		LR33	NID 057418 1.	7	NP_057418.1	
Receptor	Cystelnyl Leukotrlene CYSLT2 NP_065110.1	Cysteinyl Leukotrlene CYSLT2 Receptor	Leukotriene	Cysteinyl Leukotriene CYSLT2 NP_065110.1 Receptor	Leukotriene		Leukotriene	G Protein-Coupled	Receptor C5/2 G Protein-Coupled	Receptor C512 G Protein-Coupled	Receptor C5L2 G Protein-Coupled	Receptor C5L2 G Protein-Coupled	Receptor C5L2 G Protein-Coupled	Receptor Ls 190438 G Protein-Coupled	Receptor Ls190484	G Protein-Coupled Receptor Ls 190484	© Protein-Coupled	receptor LS190484 G Protein-Coupled	Receptor Ls 190484 G Protein-Coupled	Receptor SH120	G Protein-Coupled	27:10:10:10:10:10:10:10:10:10:10:10:10:10:
	190427	190427	190427	190427	190427	190427	190427	190437	190437	190437	190437	190437				190484 (190484	190484	100505		190595	
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KSVTTSASGSENLTLLQQE	EVDALEELSROLFLETAD	DRVGKTDPVTRGIEIT		VRLPFIKEKEKKSPVGLH		DEHNAALRIAGFPNGSLGKR	GKRPSGSLGKRPSAPFRSNV		SQPRMRETAFEEDVQLPR		GDPAIYQSLKAQNAYSRHC	PESSHSSVTVPSKKIFI SKI		GKILLNILTLGMRRKNTCQN		EEVITLVQAIRITSYMNE		CKGNGESLWQRQRLQSE	RHSRPYPSYRSTHRST	TSHTSNLSWISIRRRQE	DLEAKAPPRPQGHEAET	KLGRRPVAVDVLLLNLTASD			EFSGDISHSQGTNGTC		SRLVWILGRGGSHRRQRR		GGWQQESSMELKEQKGG		EEQRADRPAERKTSEHSQGC	MOTGPOGSVESGNHWEVESV	
1732	1733	1734		411		412	413		414	(542	543		619		970		2137	2138	2139	2140	1735	1736	3	1737		1738		1739		1740	2569))
NP_057418.1	NP_057418.1	NP_057418.1		075205		0/5205	075205		075205		CAB55314.1	CAB55314.1		CAB55314.1		CAB55314.1		AAF24978.1	AAF24978.1	AAF24978.1	AAF24978.1	NP_005295.1	NP 005205 1		NP_005295.1		NP_005295.1		NP_005295.1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	NP_005295.1	NP 005295 1	
G Protein-Coupled	Receptor SH 120 G Protein-Coupled Receptor SH 120	G Protein-Coupled	Receptor SH120	G Profein-Coupled	Receptor GPRC5B	G Protein-Coupled	G Protein-Coupled	Receptor GPRC5B	G Protein-Coupled	Receptor GPIRC5B	G Protein-Coupled Receptor GPCR150	G Profein-Coupled	Receptor GPCR150	G Protein-Coupled	Receptor GPCR150	G Protein-Coupled	Receptor GPCR150	Melanopsin	Melanopsin	Melanopsin	Melanopsin	G Protein-Coupled	Receptor GPR41 & GPR42 G Protein-Coupled	Receptor GPR41 & GPR42	G Protein-Coupled	Receptor GPR41 & GPR42	G Protein-Coupled	Receptor GPR41 & GPR42	G Protein-Coupled	Receptor GPR41 & GPR42	G Protein-Coupled	Receptor Gride & Gride G Protein-Coupled	The Control of the Co
190595	190595	190595		190599		64CD4	190599		190599	00,001	70007	190602		190602	4	190602		190623	190623	190623	190623	190627	190627		190627		190627		190627		190627	190627	
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Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sanlens	Homo sapiens	Homo saplens	Homo sapiens
VAIYAYYKKQRTKTDV	VAVTKVPSQSGVGKPCWII	CNMSKRMDIAIQVTESI	ROSVEEFPFDSEGPTEP	GHPPGSGGAESADTEARVR	HSVASALKSHRTRGHGRGDC	KGGAAVAGGRPTGASARR	CLVRREFRKALKSLLWR	RPFTATTKPEHEDQGLQ	AFPPVLDVGTYSFIREEDQC	HDRRKMKPVQFVAAVSQN	RRRLLVLDEFKMEKRISR	LRRCFSTTLYCRKSRLPRE	PLTLAGVVARRQPAGDRLC	CSRRPDERLRFAVFIGA	CKEILNRLLHRRSIHSSG	CLEEGKRRRGRATKKIST	FPFFVSGAI SPPSASAVVK	NGHAASRRLLGMDEVKGEK	KKCLRTHAPCWGTGGAPAPR	VLIMAAIHAVYGKLLLFEYK
1441	1442	1443	1444	1741	1742	1743	1744	1745	339	340	341	342	554	555	557	567	516	519	526	170
AAF61299.1	AAF61299.1	AAF61299.1	AAF61299.1	NP_057652.1	NP_057652.1	NP_057652.1	NP_057652.1	NP_057652.1	CAB82307.1	CAB82307.1	CAB82307.1	CAB82307.1	LR26	LR26	LR26	1R26	6VI	627	627 627	
Receptor GPR41 & GPR42 C-C Chemokine Receptor	C-C Chemokine Receptor	C-C Chemokine Receptor	C-C Chemokine Receptor	G Protein-Coupled December 881 00	G Protein-Coupled	Receptor sality G Protein-Coupled December 84180	G Protein-Coupled	Receptor SALPR G Protein-Coupled	Receptor SALPIX G Protein-Coupled	G Protein-Coupled	Receptor GPR85 (SREB2) G Protein-Coupled	Keceptor GPR85 (SREBZ) G Protein-Coupled	Receptor GPR85 (SRE52) G-Protein-Coupled Docortor GPP24	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	Receptor Gritzo Sreb3	Sreb3	Sreb3	CODIO
190701	190701	190701	190701	190705	190705	190705	190705	190705	190711	110011	190711	190711	190725	190725	190725	190725	190741	190741	190741	1 () (
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Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens			Homo sapiens	Homo sapiens	Homo sapiens	Homography		Homo saplens		Homo sapiens	Homo sapiens		Homo saplens	Homo saplens		Homo sapiens	Homo sanlens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
RRAPGPPSDTFVFNLALAD	GRRGRRRGDSRVVARSVR	RREPROALAGIFRDLIRSR	KGVGRRWVASNPRESRPS	KDCIESTGDYFLLCDAFGP	VENIOEI SPOTEI ONGOS		GUSGSKEVLLGEKGEKNHA	SMLLRGNPQFQRQPQWDDP	KVPSEELTTSSSHGPPPTAR	VOSCECED DONGS A CWAVY		QDTKKRSLLGTQVFFLLGT		KECKG-CSIMI-VENKAFSIMIDE	TATEIRNOVKKEMILAKR		NYRQRKSMDSKG@KTYAPS	SCSNLTVLVMRKNKINHLN		DELDLGSINKIENLYPLITKU	OI SSPSPPTCIKTI CSI R	DMLKIASMHSQQIRKMEHAG.	AGGYRSPRTPSDFKALRTVS	RESSCHIVTISSSEFDG	GVKKVLTSFLLFLSARNC	NSLLNPLIYAYWQKEVRLQ	RRAALRPPRPARGSRLRSD
. 550	551	552	553	568	27,7	, CG	0/9	571	529	530	700	535	CCL	979	260		561	565		000	546	547	548	549	1481	1482	467
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LR23	EZ31	LR23	LR23	LR32	1.030	701	11432	LR32	LR34	1034	5	LR34		LK34	LR40		LR40	LR40	9	LK40	1 647	LR47	LR47	LR47	LR47	LR47	LR48
G Protein-Coupled	Receptor H71BA62 G Protein-Coupled Receptor H7TBA62	G Protein-Coupled	Receptor H71BA62 G Protein-Coupled	Receptor H7TBA62 G Protein-Coupled	Receptor GPRC5D	Receptor GPRC5D	G Protein-Coupled Receptor GPRC5D	G Protein-Coupled	G-Protein-Coupled	Receptor GPRC5C	Receptor GPRC5C	G Protein-Coupled	Receptor GPRC5C	Perentor Coupled	G Protein-Coupled	Receptor LGR7	G Protein-Coupled Receptor I GR7	G Protein-Coupled	Receptor LGR7	G Protein-Coupled	Receptor Lerk/ GPCR Ls190748	GPCR Ls190748	GPCR Ls190748	GPCR L3190748	GPCR Ls 190748	GPCR Ls 190748	G Protein-Coupled
190742	190742	190742	190742	190743	1007/43	00.75	190/43	190743	190744		<u> </u>	190744	1001	190/44	190745		190745	190745		190/45	190748	190748	190748	190748	190748	390748	190749
2113	2114	2115	2116		21.18	2 5	2117	2120	2121	2100	7717	2123		7174	2125		2126	2127		7170	2129	2130	2131	2132	2133	2134	2135

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	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo saplens		Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	• • • • • • • • • • • • • • • • • • • •
	RPVRLALGRLSRRALPGPVR	DSRISILPPLRPRLPGGK	RPPEGPAVGPSEAPEQTPE	VVARRAALRPPRPA	PSEAPEQTPELAGGR	GPSEAPEQTPELAG	PDTNSTINI SI STRVTI AFF	VVDKNLRHRSSYFFLN	LYIPHTLFEWDFGKEIC	TQHTGVLKIVTLMVAV	VNGPMILVSESWKDEGSEC	CEPGFFSEWYILAITSFL	AYFNMNIYWSLWKRDHLSRC	CGHSFRGRLSSRRSLS	IASKMGSFSQSDSVALHQRE	IVLSFYSSATGPKSVWYRIA	IIRVITVPGKTGTVAC	SPWTNDPKERINVAVA	RIRELLQGMYKEIGIAVD	TOTSDIATNSTIPSAE		TEVPDSA@TSNTHTTSAS •	GDTAVERLNVFITMAKV	MSLAKRVMTGLWIFTI	LHFIIGFTVPMSIITV	
	468	510	ธาา	2702	2703	2704	2235	2237	2240	2242	2243	2244	2245	2246	2247	2249	2085	2086	2087	2088		481	522	523	525	
	LR48	LR48	LR48	LR48	LR48	LR48	NP 067637.2	NP_067637.2	NP_067637.2	NP_067637.2	NP_067637.2	NP_067637.2	NP_067637.2	NP_067637.2	NP_067637.2	NP_067637.2	NP_002020.1	NP_002020.1	NP_002020.1	NP 002020.1		LR14	LR14	LR14	LR14	
÷.	Receptor GPR62 G Protein-Coupled	Receptor Grivoz G Protein-Coupled	Receptor GPR62 G Protein-Coupled	Receptor GPR62 G Protein-Coupled	Receptor GPR62 G Protein-Coupled	Receptor GPR62 G Protein-Coupled	Receptor GPR62 Histamine H4 Receptor	Histamine H4 Receptor	Histamine H4 Receptor	Histamine H4 Receptor	Histamine H4 Receptor	Formyl Peptide Receptor 1	Formyl Peptide Receptor 1 (FPD)	Formyl Peptide Receptor 1	(FPRT) Formyl Peptide Receptor 1	(FPR1)	Formyl Peptide Receptor- like 2 (FPDI 2)	Formyl Peptide Receptor-	Formyl Peptide Receptor	like 2 (FPRL2) Formyl Peptide Receptor-						
	190749	190749	190749	190749	190749	190749	190774	190774	190774	190774	190774	190774	190774	190774	190774	190774	190823	190823	190823	190823		190824	190824	190824	190824	
	2136	2137	2138	2139	2140	2141	2142	2143	2144	2145	2146	2147	2148	2149	2150	2151	2152	2153	2154	2155		2156	2157	2158	2159	٠.

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	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens		Homo sapiens	Homo saplens		Homo saplens	Homo sapiens		Homo sapiens	Homo sapiens		Homo sapiens	Homo saplens	Homos capiens		Homo saplens	Homos carolons	si piche ci ibili	Homo sapiens		sueidos oution	Homo sapiens	Homos caron	suppose of their	
	DELLEAPGDLETLPRLQQHC CVASHI DGI FDVI RGI SKN	KSGDPGPSVVGLVSIPG	SKGIRKLKTESEMHTLSSS	ELSLEVQKQVDRSVTLRQNQ	EPEKOMLLHETHQGLLQDGS	KRMQKRSVTALMVLNLALAD		RPFVSQKLRTKAMARR	ASYSDIGRRLQARRER		LEGTGSEASSTRRGGS	RKALKMMLFGKIFQKDSSRC		QIGLEMKNGISGSKERKAV	RIYLIAKEQARLISDANQK		ELNFKGAEEIYYKHVHC	CVKNNWSNDVRASLYS	SAFPPADWDGAGGSVRITRG		GIVRRVRVSVKRVSVLN	DNIEFERDS//PS// PG//CD&		CEEESWAGRRIPVSLLYSG		C L'EUNINA INVOVININO	KELYRSYVRTRGVGKVPR	SOXXXIVX COO COINT II		
	1658 1659	1660	1661	1662	1663	1492		1493	1494		1495	2039		2040	2041	. (2042	2043	1569	ì	1571	1572	2	1573	1971	3	1544	15/15	3	
	NP_038475.1 NP_038475.1	NP_038475.1	NP_038475.1	NP_038475.1	NP_038475.1	NP_000743.1		NP_000743.1	NP_000743.1		NP_000743.1	LR122	•	LR122	LR122	9	UR122	LR122	NP 071332.1		NP_071332.1	NP 0713321		NP_071332.1	1 000170 GIA	147_07 1932.1	NP_073625.1	NP 0734251	-0.0000	
IIKO Z (FFIKLZ)	EMR2 Hormone Receptor EMR2 Hormone Receptor	EMR2 Hormone Receptor	EMR2 Hormone Receptor	EMR2 Hormone Receptor	EMR2 Hormone Receptor	Leukotriene B4 Receptor	BLT1	Leukotriene B4 Receptor 8171	Leukotriene B4 Receptor	BLTT	Leukotriene B4 Receptor	Trace Amine Receptor 1	(TA1)	Trace Amine Receptor 1 (TA1)	Trace Amine Receptor 1	(IAI)	Iface Amine Receptor I	Trace Amine Receptor 1	(IAI) G Protein-Coupled	Receptor 88 (GPR88)	G Protein-Coupled	receptor oo (GP1389) G Protein-Coupled	Receptor 88 (GPR88)	G Protein-Coupled	Receptor 88 (GPR88)	Receptor 88 (GPR88)	P2Y12 Platelet ADP	Receptor P2V12 Pintelet ADP	Receptor	
	190948 190948	190948	190948	190948	190948	190955		190955	190955		190955	191039		191039	191039	060101	650161	191039.	191132		191132	101132		191132	051101	707	191168	9181 191168	3	
	2160 2161	2162	2163	2164	2165	2166	:	2167	2168	:	2169	2170		2171	2172	0170	21/3	2174	2175		2176	7716		2178	07.10		2180	2181	<u>-</u> -	

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Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens
CPNSATSLSQDNRKKEQDGG	TTRPFKTSNPKNLLGAK	ANEGGIEELVVA	RKIESTASQAQSS	LVDAVIDAYMNFI	RIDSSTINLFSEEVET	NASDFPDYAAAFGNCTDE	TFLITSTNRTNRSACLD	TLTHGLQTDSCLKQKARR	RLLSISCSIENQIHEA	QQAVCSTVRCKVSGNLE	QDIAEVDHSEGCF	RKGWRLGQPILKLA	CSISINFPSFFTTVMTC	QWFLILWIWKDSDV	AFLSDNTIEVRINRTLKK	QETKNEFRNLKQIQSKC	CNNKTHWAPVRSTM	TKMAEYDLQNDVFIIPD	CQDTTSKTTEGRKELQKIV
1546	1570	6961	2316	2571	2573	1864	1865	1866	1867	1868	2749	2750	2751	2752	2575	2576	2577	2581	1665
NP_073625.1	NP_073625.1	LR88	LR88	LR88	LR88	IP_13092	IP_13092	IP_13092	IP_13092	IP_13092	AAK91805.1	AAK91805.1	AAK91805.1	AAK91805.1	ENSP00000199719	ENSP00000199719	ENSP00000199719	ENSP00000199719	AAK15076.1
P2V12 Platelet ADP	keceptor P2Y12 Platelet ADP Receptor	Trace Amine Receptor 3	Trace Amine Receptor 3	Trace Amine Receptor 3	(1A3) Trace Amine Receptor 3	(173) G Protein-Coupled Receptor GPR80	G Protein-Coupled Recentor GPR80	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	Receptor GPR80 MrgX2 G Protein-Coupled	MrgX2 G Protein-Coupled	MgX2 G Protein-Coupled Recentor	MrgX2 G Protein-Coupled	Receptor G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	receptor 13191222 EGF-Like Module-Containing AAK15076.1
191168	191168	191193	191193	191193	191193	191196	961161	191196	961161	961161	191218	191218	191218	191218	191222	191222	191222	191222	193511
2182	2183	2184	2185	2186	2187	2188	2189	2190	2191	2192	2193	2194	2195	2196	2197	2198	2199	2200	2201

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Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens		Homo sapiens	Homo sapiens	Homos canciens		Homo sapiens	Homo soniens		Homo sapiens		Homo sapiens		Homo saplens		Homo sapiens			Homo sapiens		Homo sapiens			Homo sapiens	•	
RDVESKVLETALKDPEQK	KIQNDSVAIETQAITDNC	CSEERKTFNLNVQMNSMDIR	EEMDKKDQVYLNSQVVSAA		SKSVILIFCHVKMIPSIK	CLLLPTAVIVFSYVKIIAK	A LITA/A/P (C)GIPCOG		CQTGGLKATKKKSLEG	PI HTVTTVPKSSAVI F		PTAVIVESYVKIIAKV		KLAGI?LI?EVIGHIDHYFSQD		CALQTWGSERRLGLDTSKD		RGRRQSARNSRGPPEQPNE			INDIAGPPECPINEELG		A BOVRED VRPHTVVLR			QLDQVPSRHPSRE		-
1666	1667	1668	6991	0	0/01	2142	77/10	<u> </u>	2145	2146	<u>1</u>	2620		194/		1948		2734			2/35		2736	; ;	:	2742		
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AAK15076.1	AAK15076.1	AAK15076.1	AAK15076.1		AAK150/6.1	CAC21687.1	CAC21687 1		CAC21687.1	CAC21687 1	(1001.1	CAC21687.1		NP_001398.1		NP_001398.1		NP_001398.1	· .		NP_001398.1		NP 001398.1			NP_001398.1	-	
MR3 aining	EGF-Like Module-Containing	FGF-Like Module-Containing	Mucin-Like Receptor EMR3 EGF-Like Module-Containing	Mucin-Like Receptor EMR3	EGF-Like Module-Containing Mucin-Like Receptor EMR3	G Protein-Coupled	Receptor dJ402H5.1 G Protein-Compled	Receptor dJ402H5.1	G Protein-Coupled	Receptor dJ402H5.1 G Protein-Coupled	Receptor dJ402H5.1	G Profein-Coupled	Receptor dJ402H5.1	Cadherin EGF LAG Seven-	Pass G-Type Receptor 3 (CELSR3)	Cadherin EGF LAG Seven-	Pass G-Type Receptor 3	EGF LAG Sev	Pass G-Type Receptor 3	(CELSK3)	Cadheiln EGF LAG Seven-	Pds (5-1ype keceptor 3	Cadherin EGF LAG Seven-	Pass G-Type Receptor 3	(CELSR3)	Cadherin EGF LAG Seven-	Pass G-Type Receptor 3	(Crumic)
193511	193511	193511	193511		19391	193516	103516	2	193516	103516	2	193516		193524	٠	193524		193524			193524		193524	.*		193524		
2202	2203	2204	2205		9077	2207	220R		2209	2210	2	2211	. 6	2212		2213		2214		,	2712		2216			2217		

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Homo saplens			Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens		Homo sapiens	· · ·	Homo sapiens		Homo saplens		Homo saplens	-	Homo sapiens	-	Homo saplens		Homo sapiens		Homo saplens		Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	·	Homo sapiens	٠	Homo sapiens		Homo sapiens	
LDSLSRSSNSREQLDQV			REEHHFMVDARNRSYPLYSC	PGPAPGGEEAADPRASRR	CPRPSGSHKEAYSERPGGLL	PSSGAPRPGRLPLRNGRVA	FLGKNDDIKTKKELIVN		QVTYRDSKEKRDLRNFLK		CERTKIWGTFKINERFIND		SKYANGIEIQLKKAYER		CIVVFIVRTERSLHAP		KILALFWFDSREISFEAC		CVHQDVMKLAYADTLP		RFGNSLHPIVRVVMGD		KTKQIRTRVLAMFKISC		KTDENEGDGSASVDMVFSP	KKDYQYPKSLDILSNVGC	KNLQTSDGDINNIDFDNN	SONGNINPOWELDYROEKIC	RPRLRVKMYNFLRSLPTLHE	CNPSVPKQRVMKLTKM	•	RLTRWRTRYKTIRIŃLG		KDGVESCAFDLTSPDDVL		LSGNFQKRLPQIQRRATE	
2744			1903	1904	1905	1906	2018		2019		2020		2021		2022		2023		2024		2027		2028	1 1 1	1855	1856	1857	1858	1859	1845		1846		1847	•	1848	
NP_001398.1			NP_071429.1	NP_071429.1	NP_071429.1	NP_071429.1	NP_079324.1	Į.	NP_079324.1		NP_079324.1		NP_079324.1		NP_110401.1	÷	NP_110401.1	-	NP_110401.1		NP_110401.1		NP_110401.1		LR77	LR77	LR77	LR77	LR77	AAK32193.1		AAK32193.1		AAK32193.1		AAK32193.1	
Cadherin EGF LAG Seven-	Pass G-Type Receptor 3	(CELSR3)	Neuropeptide FF 1 Receptor	G Protein-Coupled	Receptor FLJ22684	G Protein-Coupled	Receptor FLJ22684	G Protein-Coupled	Receptor FLJ22684	G Protein-Coupled	Receptor FL/22684	Olfactory Receptor, Family	51, Subfamily E, Member 2	Olfactory Receptor, Family	51, Subfamily E, Member 2	Olfactory Receptor, Family	51, Subfamily E, Member 2	Olfactory Receptor, Family	51, Subfamily E, Member 2	Olfactory Receptor, Family	31, Subjecting E. Member 2	FLJ 14454	FLJ14454	FU14454	FLJ14454	FL)14454	G Protein-Coupled	Receptor SLI/MCH2	G Protein-Coupled	Receptor SLT/MCH2	G Protein-Coupled	Receptor SLI/MCH2	G Protein-Coupled	Receptor act/inichz			
193524			193914	193914	193914	193914	194319		194319	•	194319		194319		194431		194431		194431		194431		194431		194/43	194743	194743	194743	194743	194745		194745		194745		194/45	
2218			2219	2220	222	2222	2223		2224		2225		2226		2227	-	2228		2229		2230		2231		7577	2233	2234	2235	2236	2237		2238		2239	. (2240	

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Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens	Homo sapiens
TIIRSRKTVPDIVIC	RRATEKEINNMGNTLKSHF	CRIEGDTISQVMPPLLIVA	RRHWAFGDIPCRVGLFTL	CESFIMESANGWHDIM	CSFKIVWSLRRRQQLARQAR	RRRAGLARGARMKKATR	TVPSSACDPSVHGALH	CSLKPKQPGHSKTQRPEEM	CISVANSFQSQSDGQWD	RTRKOHSEATNSSNRVFVYC	RVISQISADNYKIHGDPSA	TSSSARTSNAKPFHSD	NGTRPGMASTKLSPWD	LGIAWDRRLRSPPAGC	GERYMAVLRPLQPPGS	CRDEPSALARALTWRQAR	AAQRCLQGLWGRASRD	RDSPGPSIAYHPSSQSSVD	ALFSRIHLDWKVLF
1849	1907	2089	2090	2091	2092	2093	2094	2095	2096	2034	2035	2036	2037	1933	1934	1935	1936	1937	2748
AAK32193.1	AAK32193.1	AAK29071.1	AAK29071.1	AAK29071.1	AAK29071.1	AAK29071.1	AAK29071.1	AAK29071.1	AAK29071.1	CAB82385.1	CAB82385,1	CAB82385.1	CAB82385.1	LR84	LR84	LR84	LR84	LR84	AAK91806.1
G Protein-Coupled	G Protein-Coupled Deceptor St 7/4/CH2	Chemokine Receptor	Chemokine Receptor	Chemokine Receptor	Chemokine Receptor	Chemokine Receptor	FKSG80/GPR81 Chemokine Receptor	Chemokine Receptor	Chemokine Receptor	rnseau/erkal G Protein-Coupled Recentor I s104757	G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	Receptor LS 194/5/ G Protein-Coupled	Receptor LS194858 G Protein-Coupled	Receptor LS194858 G Protein-Coupled	G Protein-Coupled	G Protein-Coupled	Mecepior La 194838 MrgX3 G Protein-Coupled
194745	194745	194756	194756	194756	194756	194756	194756	194756	194756	194757	194757	194757	194757	194858	194858	194858	194858	194858	194878
2241	2242	2243	2244	2245	2246	2247	2248	2249	2250	2251	2252	2253	2254	2255	2256	2257	2258	2259	2260

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	Homo sapiens	Homo saplens	Homo saplens			Homo sapiens	Homo saplens	Homo sapiens	Homo saplens	.•	Homo sapiens	Homo sapiens	• :	Homo saplens		Homo saplens		Homo sapiens	Homos omolens		Homo saplens		nomo sapiens	Homo saplens	-	Homo sapiens	مروامی مسمل		Homo saplens	
	CIAFKDIMPFSAQVGDER	KAFEEAYARADKKAPRPC	ETKIQWHGKDNQVPKSVC	7 × 014×2400 10×10 × 30	Catcondirect	SDYDMPLDEDEDVTNS	NPHGAHATSFPFNFSY	ERALPRIYMASVYNTRHVC	CAKMQNAEAADATLVF		DRDTGRLEPSAHRLLVATVC	RYMNGSFPSKLQRLMKKLPC		CARAAGDAPLRSLEQANRTR		VISYSKILOTTKASRKRL		TVSLAYSRSHQIRVSQQD	CHWEPEKGAII TOTSVKRND		TYGRDNGQLLGERVARRDIC		GEILPILGPINGINMISEEKOK	RTSQSYTCNQECDNCLNAT .		RPQSHPRTDPDDPKITIVSC	1/3 (1/4 1/2) V (1/4 4/4)	VAINTENTOON	KVIVTGQVLKNSSA	
	1991	1992	1993	200	1444	2011	2014	1986	1987		1988	1989	- -	2003		2004		2005	2006		2007	0000	2008	2009		2010		7177	2313	
	ENSP00000198236	ENSP00000198236	ENSP00000198236	7 COOCOCCANT	ENSPUCIONISSES	LR114	LR114	LR112	LR112		LR112	LR112		LR116		LR116		LR116	. 91101		LR117	!	חלוו/	LR117		LR117	. 0701774	AAK/ 1243. I	AAK71243.1	
Receptor	G Protein-Coupled Receptor GPCR83	G Protein-Coupled	Receptor GPCRB3 G Protein-Coupled	Receptor GPCRB3	G Protein Coupled Receptor GPCRB3	WO0034334-hFB41A	WO0034334-hFB41A	G Protein-Coupled	G Protein-Coupled	Receptor MGC7035	G Protein-Coupled	Receptor MGC/033 G Protein-Coupled	Receptor MGC7035	G Protein-Coupled	Receptor 14273	G Protein-Coupled	Receptor 14273	G Protein-Coupled	Receptor 142/3 G Protein-Coupled		G Protein-coupled Receptor LR117	Gpc104	G Protein-coupled Receptor UTITY	G Protein-coupled Receptor LR117	Gpcrb4	G Protein-coupled Receptor LR117	Sports Programme Transfer of	irace Amine keceptor 4 (TA4)	Trace Amine Receptor 4	(IA4)
	194903	194903	194903	10,000	194903	194904	194904	194905	194905		194905	194905		194907	•	194907	~ !	194907	10/0/7		194908		194908	194908		194908	2000	19493/	194957	
	2261	2262	2263	3	4077	2265	2266	2267	2268		2269	2270		2271		2272		2273	7,777	r }	2275		2276	2277		2278	0	6/77	2280	

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Homo sapiens	Homo sapiens	Homo sapiens	Homo saplens	Homo sapiens	Homo sapiens	Homo saplens	Homo saplens	Homo saplens	Homo saplens	Homo sapiens	Homo sapiens
MSSNSSILVAVQLC	IAKQQAIKIETISSKV	MISNESQPVVQLC	KULSGDVLKAS	SGDVLKASSSTISLFLE	QDKPEVDKGEGQLPEESL	CINISHURKILVS	MDPTVPVFGTKL	RYATLMQKDSSQETT	KIFYGHLLKKFRQPNF	YSVIEATEGEESLC	CTSIMEKDLTYSSVKR
2318	2307	2314	2319	2570	2727	2728	2729	2706	2707	2708	2715
AAK71243.1	AAK71244.1	AAK71244.1	AAK71244.1	AAK71244.1	AAK91807.1	AAK91807.1	AAK91807.1	AAL26482	AA126482	AAL26482	AAL26482
	Trace Amine Receptor 5	Irace Amine Receptor 5	Irace Amine Receptor 5	Irace Amine Receptor 5	MrgX4 G Protein-Coupled AAK91807.1	MrgX4 G Protein-Coupled	peldr	G Protein-Coupled	G Protein-Coupled	Receptor GPR82 G Protein-Coupled	receptor GPR82 G Protein-Coupled Receptor GPR82
2281 194957	194958	194958	194958	194958	194989	194989	194989	195015	195015	195015	195015
2281	2282	2283	2284	2285	2286	2287	2288	2289	2290	2291	2292

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SEQ ID NO:	LS_ID	Gene	Antibody Company Name
1	127	5-HT1A Receptor	Chemicon
1	127	5-HT1A Receptor	Research Diagnostics
ī	127	5-HT1A Receptor	Santa Cruz
3	128	5-HT1B Receptor	Chemicon
3	128	5-HT1B Receptor	Research Diagnostics
3	128	5-HT1B Receptor	Santa Cruz
5	129	5-HT1D Receptor	Research Diagnostics
5	129	5-HT1D Receptor	Santa Cruz
11	132	5-HT2A Receptor	Calbiochem
11	132	5-HT2A Receptor	Research Diagnostics
13	133	5-HT2B Receptor	Research Diagnostics
15	134	5-HT2D Receptor	Research Diagnostics
	134	5-HT2C Receptor	Santa Cruz
15		5-HT7 Receptor	Calbiochem
21	139		Alpha Diagnostic Int.
23	272	Adenosine Al Receptor	Calbiochem
23	272	Adenosine A1 Receptor	Santa Cruz
23	272	Adenosine A1 Receptor	
25	273	Adenosine A2a Receptor	Alpha Diagnostic Int.
25	273	Adenosine A2a Receptor	Calbiochem
25	273	Adenosine A2a Receptor	Chemicon
25	273	_	Santa Cruz
27	274	Adenosine A2b Receptor	Alpha Diagnostic Int.
27	274	Adenosine A2b Receptor	Chemicon
27	274	Adenosine A2b Receptor	Santa Cruz
29	275	Adenosine A3 Receptor	Alpha Diagnostic Int.
29	275	Adenosine A3 Receptor	Santa Cruz
31	309	Melanocortin 2 Receptor	Alpha Diagnostic Int.
		(adrenocorticotropic hormone) (MC2R)	
31	309	Melanocortin 2 Receptor	Chemicon
		(adrenocorticotropic hormone)	
		(MC2R)	
31	309	Melanocortin 2 Receptor	Research Diagnostics
J. -		(adrenocorticotropic hormone)	<u> </u>
		(MC2R)	
31	309	Melanocortin 2 Receptor	Santa Cruz
J1 .	307	(adrenocorticotropic hormone)	
	•	(MC2R)	
25	377	Alpha 1b-adrenoceptor	Research Diagnostics
35			Santa Cruz
35	377	Alpha 1b-adrenoceptor	Research Diagnostics
37	379	-	Santa Cruz
37	379	Alpha 1c-adrenoceptor	
39	387	Alpha 2a-adrenoceptor	Calbiochem
39	387	Alpha 2a-adrenoceptor	Santa Cruz
41	388	Alpha 2b-adrenoceptor	Research Diagnostics
41	388	Alpha 2b-adrenoceptor	Santa Cruz
43	389	Alpha 2c-adrenoceptor	Research Diagnostics
43	389	Alpha 2c-adrenoceptor	Santa Cruz
.45	599	Bradykinin B1 Receptor	Research Diagnostics
49	635	Beta-1 adrenoceptor	Calbiochem
49	635	Beta-1 adrenoceptor	Research Diagnostics
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192	3226	Muscarinic acetylcholine Receptor M4	Santa Cruz	
194	3227	Muscarinic Acetylcholine Receptor M5	Biogenesis	
194	3227	Muscarinic Acetylcholine Receptor M5	Santa Cruz	
200	3404	Neuropeptide Y Receptor Type 2	Biogenesis	
202	3405	Neuropeptide Y Receptor Type	Biogenesis	•
206	2400		Santa Cruz	
206	3408	Neurotensin Receptor Type 1	· ·	
208	3452	Opiate Receptor-Like 1 (OPRL1)	Santa Cruz	· · · · · · · · · · · · · · · · · · · ·
214	3582	Oxytocin Receptor	Santa Cruz	
216	3589	Purinergic Receptor P2Y, G- protein coupled, 2 (P2RY2)	Chemicon	
216	3589	Purinergic Receptor P2Y, G- protein coupled, 2 (P2RY2)	Zymed	
218	3595	Purinergic Receptor P2Y1	Chemicon	1.5
218	3595	Purinergic Receptor P2Y1	Zymed	
228	3640	Parathyroid Hormone Receptor		
	551,5	1 (PTHR1)		
228	3640	Parathyroid Hormone Receptor 1 (PTHR1)	Lab Vision Corporatio	n/NeoMarkers
228	3640	Parathyroid Hormone Receptor 1 (PTHR1)	Santa Cruz	
236	3846	Sphingolipid Receptor Edg1	Exalpha Biologicals	
238	3847	Sphingolipid Receptor Edg3	Exalpha Biologicals	
240	3848	C-C Chemokine Receptor 9	Research Diagnostics	
248	3852	CX3C Chemokine Fractalkine	Chemicon	
240	3032	Receptor 1	Chemicon	
248	3852		Chemokine.com	
•	'	Receptor 1		
248	3852	CX3C Chemokine Fractalkine Receptor 1	, Artist	
250	3853	G Protein-Coupled Receptor GPR15	Santa Cruz	
264	3860	G Protein-Coupled Receptor SLC/MCH1	Alpha Diagnostic Int.	
264	3860	G Protein-Coupled Receptor SLC/MCH1	Santa Cruz	3434
295	3927	Prostaglandin E Receptor EP4	Cayman	
299	4051	Proteinase-Activated Receptor	Research Diagnostics	
	.00.	2		
299	4051	Proteinase-Activated Receptor 2	Santa Cruz	
301	4052	Proteinase-Activated Receptor	Research Diagnostics	
301	4052	Proteinase-Activated Receptor	Santa Cruz	
205	1251	3. Phodonoin	Biocarta	
305	4254	Rhodopsin		
305	4254	Rhodopsin	DPC Biermann/Acris	
311	4480	Somatostatin Receptor Type 1	Santa Cruz	

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49	635	Beta-I adrenoceptor	Santa Cruz
51	640	Beta-2 adrenoceptor	Research Diagnostics
51	. 640	Beta-2 adrenoceptor	Santa Cruz
53	643	Beta-3 adrenoceptor	Alpha Diagnostic Int.
53	643	Beta-3 adrenoceptor	Chemicon
53	643	Beta-3 adrenoceptor	Research Diagnostics
53	643	Beta-3 adrenoceptor	Santa Cruz
57	692	Bombesin Receptor Subtype-3	Alpha Diagnostic Int.
57	692	Bombesin Receptor Subtype-3	Chemicon
59	729	CXC Chemokine Receptor 5	Research Diagnostics
59	729	CXC Chemokine Receptor 5	Santa Cruz
61	735	C-C Chemokine Receptor 1	Calbiochem
61	735	C-C Chemokine Receptor 1	Capralogics
61	735	C-C Chemokine Receptor 1	Chemicon
61	735	C-C Chemokine Receptor 1	Research Diagnostics
61	735	C-C Chemokine Receptor 1	Santa Cruz
63	737	C-C Chemokine Receptor 3	Research Diagnostics
63	737	C-C Chemokine Receptor 3	Santa Cruz
65	738	C-C Chemokine Receptor 4	Capralogics
65	738	C-C Chemokine Receptor 4	Research Diagnostics
65	738	C-C Chemokine Receptor 4	Santa Cruz
67	741	C-C Chemokine Receptor 7	Research Diagnostics
67	741	C-C Chemokine Receptor 7	Santa Cruz
69	742	C-C Chemokine Receptor 8	Chemicon
70	742	C-C Chemokine Receptor 8	Chemicon
71	742	C-C Chemokine Receptor 8	Chemicon
73 ·	· 752	CXC Chemokine Receptor 3	Research Diagnostics
73	752	CXC Chemokine Receptor 3	Santa Cruz
73	752	CXC Chemokine Receptor 3	Zymed
75	753	CXC Chemokine Receptor 4	Biosource
75	753	CXC Chemokine Receptor 4	Calbiochem
75	753	CXC Chemokine Receptor 4	Capralogics
75	753	CXC Chemokine Receptor 4	Chemicon
75	753	CXC Chemokine Receptor 4	eBioscience
75	753	CXC Chemokine Receptor 4	Research Diagnostics
75	753	CXC Chemokine Receptor 4	Santa Cruz
77	755	Complement Component 3a Receptor 1	Chemokine.com
79	758	Complement Component 5a Receptor 1	Santa Cruz
83	832	Cannabinoid Receptor 1	Alpha Diagnostic Int.
83	832	Cannabinoid Receptor 1	Biosource
83	832	Cannabinoid Receptor 1	Calbiochem
83	832	Cannabinoid Receptor 1	Cayman
83	832	Cannabinoid Receptor 1	Chemicon
83	832	Cannabinoid Receptor 1	Santa Cruz
85	833	Cannabinoid Receptor 2	Alpha Diagnostic Int.
85	833	Cannabinoid Receptor 2	Calbiochem
85	833	Cannabinoid Receptor 2	Cayman
85 ·	833	Cannabinoid Receptor 2	Chemicon
85	833	Cannabinoid Receptor 2	Santa Cruz
97	1240	Dopamine Receptor D1	Alpha Diagnostic Int.
97	1240	Dopamine Receptor D1	Biogenesis
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97	1240	Dopamine Receptor D1	Calbiochem
97	1240	Dopamine Receptor D1	Chemicon
97 .	1240	Dopamine Receptor D1	FabGennix through Abcam
97	1240	Dopamine Receptor D1	Research Diagnostics
97	1240	Dopamine Receptor D1	Santa Cruz
99	1241	Dopamine Receptor D5	Alpha Diagnostic Int.
99	1241	Dopamine Receptor D5	Biogenesis
99	1241	Dopamine Receptor D5	Calbiochem
99	1241	Dopamine Receptor D5	Chemicon
99	1241	Dopamine Receptor D5	Santa Cruz
101	1242	Dopamine Receptor D2	Alpha Diagnostic Int.
101	1242	Dopamine Receptor D2	Biogenesis
101	1242	Dopamine Receptor D2	Calbiochem
101	1242	Dopamine Receptor D2	Chemicon
101	1242	Dopamine Receptor D2	DPC Biermann/Acris
101	1242	Dopamine Receptor D2	FabGennix through Abcam
101	1242	Dopamine Receptor D2	Research Diagnostics
101	1242	Dopamine Receptor D2	Santa Cruz
103	1243	Dopamine Receptor D3	Alpha Diagnostic Int.
103	1243	Dopamine Receptor D3	Biogenesis
103	1243	Dopamine Receptor D3	Calbiochem
103	1243	Dopamine Receptor D3	Chemicon
103	1243	Dopamine Receptor D3	Research Diagnostics
103	1243	Dopamine Receptor D3	Santa Cruz
103	1243	Dopamine Receptor D3	Zymed
105	1244	Dopamine Receptor D4	Alpha Diagnostic Int.
105	1244	Dopamine Receptor D4	Biogenesis
105	1244	Dopamine Receptor D4	Calbiochem
105	1244	Dopamine Receptor D4	Chemicon
105	1244	Dopamine Receptor D4	DPC Biermann/Acris
105	1244	Dopamine Receptor D4	Santa Cruz
107	1267	Opioid Receptor, delta 1 (OPRD1)	Biosource
107	1267	Opioid Receptor, delta 1 (OPRD1)	Calbiochem
107	1267	Opioid Receptor, delta 1 (OPRD1)	DPC Biermann/Acris
107	1267	Opioid Receptor, delta 1 (OPRD1)	Santa Cruz
113	1486	Endothelin B Receptor	Biogenesis
113	1486	Endothelin B Receptor	Capralogics
113	1486	Endothelin B Receptor	DPC Biermann/Acris
113	1486	Endothelin B Receptor	Fitgerald Industries Int.
113	1486	Endothelin B Receptor	Research Diagnostics
115	1488	Endothelin A Receptor	Biogenesis
115	1488	Endothelin A Receptor	Capralogics
115	1488	Endothelin A Receptor	DPC Biermann/Acris
115	1488	Endothelin A Receptor	Fitgerald Industries Int.
115	1488	Endothelin A Receptor	Research Diagnostics
117	1598	Calcium-Sensing Receptor	Chemicon
- *		(CASR)	
117	1598	Calcium-Sensing Receptor	DPC Biermann/Acris

1681 Follicle Stimulating Hormone Receptor Receptor	WO 02/	061087	443/448		PCT/US01/50107
1681	121	1681	Follicle Stimulating Hormone	Biogenesis	
121	121	1681	Follicle Stimulating Hormone	DPC Biermann/Acris	
1762 Galanin Receptor GalR1 Alpha Diagnostic Int.	121	1681	Follicle Stimulating Hormone	Santa Cruz	· · · · · · · · · · · · · · · · · · ·
1925 Gonadotropin-Releasing Hormone Receptor	125	1762		Alpha Diagnostic Int	
Hormone Receptor 135 1925 Gonadotropin-Releasing Hormone Receptor 136 1925 Gonadotropin-Releasing Hormone Receptor 137 1925 Gonadotropin-Releasing Hormone Receptor 138 1925 Gonadotropin-Releasing Hormone Receptor 139 1951 Growth Hormone Santa Cruz 139 1951 Growth Hormone Santa Cruz 143 2120 Histamine H1 Receptor Alpha Diagnostic Int. 145 2121 Histamine H2 Receptor Chemicon 147 2783 Opioid Receptor, kappa 1 (OPRK1) 147 2783 Opioid Receptor, kappa 1 (OPRK1) 147 2783 Opioid Receptor, kappa 1 (OPRK1) 148 21976 Lysophosphatidic Acid Receptor Edg2 155 3057 Melanocortin 3 Receptor (MC3R) 155 3057 Melanocortin 3 Receptor (MC3R) 157 3058 Melanocortin 4 Receptor (MC4R) 157 3058 Melanocortin 4 Receptor (MC4R) 159 3059 Melanocortin 5 Receptor (MC5R)					
135 1925 Gonadotropin-Releasing Hormone Receptor 136 1925 Gonadotropin-Releasing Hormone Receptor 137 1925 Gonadotropin-Releasing Hormone Receptor 138 1925 Gonadotropin-Releasing Hormone Receptor 139 1951 Growth Hormone Secretagogue Receptor 139 1951 Growth Hormone Secretagogue Receptor 143 2120 Histamine H1 Receptor Alpha Diagnostic Int. 145 2121 Histamine H2 Receptor Alpha Diagnostic Int. 146 2121 Histamine H2 Receptor Alpha Diagnostic Int. 147 2783 Opioid Receptor, kappa 1 (OPRK1) 151 2976 Lysophosphatidic Acid Receptor Edg2 155 3057 Melanocortin 3 Receptor (MC3R) 155 3057 Melanocortin 3 Receptor (MC3R) 155 3057 Melanocortin 3 Receptor (MC3R) 157 3058 Melanocortin 4 Receptor (MC4R) 157 3058 Melanocortin 4 Receptor (MC4R) 157 3058 Melanocortin 4 Receptor (MC4R) 159 3059 Melanocortin 5 Receptor (MCSR)	100,	1,20			
1925 Gonadotropin-Releasing Hormone Receptor	135	1925	Gonadotropin-Releasing	Lab Vision Corporatio	n/NeoMarkers
1925 Gonadotropin-Releasing Hormone Receptor Growth Hormone Santa Cruz	135	1925	Gonadotropin-Releasing	Research Diagnostics	
199	135	1925	Gonadotropin-Releasing	Santa Cruz	
Secretagogue Receptor 143 2120	130	1051		Santa Cruz	
143 2120	139	1931		Santa Cruz	
143 2120	143	2120		Alpha Diagnostic Int.	
145 2121 Histamine H2 Receptor Chemicon 147 2783 Opioid Receptor, kappa 1 (OPRK1) 151 2976 Lysophosphatidic Acid Receptor Edg2 155 3057 Melanocortin 3 Receptor (MC3R) 155 3057 Melanocortin 3 Receptor (MC3R) 155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Chemicon (MC3R) 157 3058 Melanocortin 4 Receptor Alpha Diagnostic Int. (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 159 3059 Melanocortin 5 Receptor Chemicon Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics					
145 2121				Alpha Diagnostic Int.	
(OPRK1) 147 2783 Opioid Receptor, kappa 1 (OPRK1) 147 2783 Opioid Receptor, kappa 1 (OPRK1) 147 2783 Opioid Receptor, kappa 1 (OPRK1) 150 Opioid Receptor, kappa 1 (OPRK1) 151 2976 Lysophosphatidic Acid Receptor Edg2 155 3057 Melanocortin 3 Receptor (MC3R) 157 3058 Melanocortin 4 Receptor (MC4R) 159 3059 Melanocortin 5 Receptor (MC5R)	145	2121			
147 2783	147	2783		Biosource	
147 2783 Opioid Receptor, kappa 1 (OPRK1) 147 2783 Opioid Receptor, kappa 1 Santa Cruz (OPRK1) 151 2976 Lysophosphatidic Acid Receptor Edg2 155 3057 Melanocortin 3 Receptor Alpha Diagnostic Int. (MC3R) 155 3057 Melanocortin 3 Receptor Chemicon (MC3R) 155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Santa Cruz (MC3R) 157 3058 Melanocortin 4 Receptor Alpha Diagnostic Int. (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 159 3059 Melanocortin 5 Receptor Chemicon Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon Chemicon Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	147	2783	Opioid Receptor, kappa 1	Calbiochem	4
147 2783 Opioid Receptor, kappa 1 (OPRK1) 151 2976 Lysophosphatidic Acid Exalpha Biologicals Receptor Edg2 155 3057 Melanocortin 3 Receptor (MC3R) 155 3057 Melanocortin 3 Receptor (Chemicon (MC3R) 155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Santa Cruz (MC3R) 157 3058 Melanocortin 4 Receptor Alpha Diagnostic Int. (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	147	2783	Opioid Receptor, kappa 1	DPC Biermann/Acris	
151 2976 Lysophosphatidic Acid Receptor Edg2 155 3057 Melanocortin 3 Receptor Chemicon (MC3R) 155 3057 Melanocortin 3 Receptor Chemicon (MC3R) 155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Santa Cruz (MC3R) 157 3058 Melanocortin 4 Receptor Alpha Diagnostic Int. (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Santa Cruz (MC4R) 157 3058 Melanocortin 4 Receptor Alpha Diagnostics (MC4R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	147	2783	Opioid Receptor, kappa 1	Santa Cruz	
155 3057 Melanocortin 3 Receptor (MC3R) 155 3057 Melanocortin 3 Receptor (MC3R) 155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Santa Cruz (MC3R) 157 3058 Melanocortin 4 Receptor Alpha Diagnostic Int. (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 159 3059 Melanocortin 5 Receptor Alpha Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	151	2976	Lysophosphatidic Acid	Exalpha Biologicals	
155 3057 Melanocortin 3 Receptor Chemicon (MC3R) 155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Santa Cruz (MC3R) 157 3058 Melanocortin 4 Receptor Alpha Diagnostic Int. (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Santa Cruz (MC4R) 159 3059 Melanocortin 5 Receptor Alpha Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	155	3057	Melanocortin 3 Receptor	Alpha Diagnostic Int.	
155 3057 Melanocortin 3 Receptor Research Diagnostics (MC3R) 155 3057 Melanocortin 3 Receptor Santa Cruz (MC3R) 157 3058 Melanocortin 4 Receptor Alpha Diagnostic Int. (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Santa Cruz (MC4R) 159 3059 Melanocortin 5 Receptor Alpha Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	155	3057	Melanocortin 3 Receptor	Chemicon	
155 3057 Melanocortin 3 Receptor Santa Cruz (MC3R) 157 3058 Melanocortin 4 Receptor Alpha Diagnostic Int. (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Santa Cruz (MC4R) 159 3059 Melanocortin 5 Receptor Alpha Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	155	3057	Melanocortin 3 Receptor	Research Diagnostics	
157 3058 Melanocortin 4 Receptor Alpha Diagnostic Int. (MC4R) 157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Santa Cruz (MC4R) 159 3059 Melanocortin 5 Receptor Alpha Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	155	3057	Melanocortin 3 Receptor	Santa Cruz	
157 3058 Melanocortin 4 Receptor Chemicon (MC4R) 157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Santa Cruz (MC4R) 159 3059 Melanocortin 5 Receptor Alpha Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	157	3058	Melanocortin 4 Receptor	Alpha Diagnostic Int.	
157 3058 Melanocortin 4 Receptor Research Diagnostics (MC4R) 157 3058 Melanocortin 4 Receptor Santa Cruz (MC4R) 159 3059 Melanocortin 5 Receptor Alpha Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	157	3058	Melanocortin 4 Receptor	Chemicon	
157 3058 Melanocortin 4 Receptor Santa Cruz (MC4R) 159 3059 Melanocortin 5 Receptor Alpha Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	157	3058	Melanocortin 4 Receptor	Research Diagnostics	
159 3059 Melanocortin 5 Receptor Alpha Diagnostic Int. (MC5R) 159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	157 .	3058	Melanocortin 4 Receptor	Santa Cruz	
159 3059 Melanocortin 5 Receptor Chemicon (MC5R) 159 3059 Melanocortin 5 Receptor Research Diagnostics	159	3059	Melanocortin 5 Receptor	Alpha Diagnostic Int.	
159 3059 Melanocortin 5 Receptor Research Diagnostics	159	3059	Melanocortin 5 Receptor	Chemicon	
	159	3059	Melanocortin 5 Receptor	Research Diagnostics	

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159	3059	Melanocortin 5 Receptor (MC5R)	Santa Cruz	
161	3061	Melanocortin 1 Receptor (MC1R)	Alpha Diagnostic Int.	
161	3061	Melanocortin 1 Receptor (MC1R)	Chemicon	
161	3061	Melanocortin 1 Receptor (MC1R)	Research Diagnostics	
161	3061	Melanocortin 1 Receptor (MC1R)	Santa Cruz	•
169	3093	Metabotropic Glutamate Receptor 1	Chemicon	
171	3094	Metabotropic Glutamate Receptor 2	Chemicon	
173	3095	Metabotropic Glutamate Receptor 3	Chemicon	
175	3096	Metabotropic Glutamate Receptor 4	Zymed	· ·
177	3097	Metabotropic Glutamate Receptor 5	Chemicon	
183	3100	Metabotropic Glutamate Receptor 8	Chemicon	
185	3212	Opioid mu-type Receptor	Biosource	+ 5
185	3212	Opioid mu-type Receptor	Calbiochem	
185	3212	Opioid mu-type Receptor	Chemicon	
185	3212	Opioid mu-type Receptor	DPC Biermann/Acris	
185	3212	Opioid mu-type Receptor	Santa Cruz	
187	3223	Muscarinic acetylcholine Receptor M1	Biogenesis	
187	3223	Muscarinic acetylcholine Receptor M1	Calbiochem	
187	3223	Muscarinic acetylcholine Receptor M1	Chemicon	
187	3223	Muscarinic acetylcholine Receptor M1	Santa Cruz	·
189	3224	Muscarinic acetylcholine Receptor M2	Biogenesis	•
189	3224	Muscarinic acetylcholine Receptor M2	Calbiochem	
189	3224	Muscarinic acetylcholine Receptor M2	Chemicon	
189	3224	Muscarinic acetylcholine Receptor M2	Santa Cruz	
191	3226	Muscarinic acetylcholine Receptor M4	Biogenesis	
192	3226	Muscarinic acetylcholine Receptor M4	Biogenesis	
191	3226	Muscarinic acetylcholine Receptor M4	Chemicon	
192	3226	Muscarinic acetylcholine Receptor M4	Chemicon	
191	3226	Muscarinic acetylcholine Receptor M4	Santa Cruz	
		•	,	•

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313	4481	Somatostatin Receptor Type 2	Biogenesis
313	4481	Somatostatin Receptor Type 2	Santa Cruz
315	4482	Somatostatin Receptor Type 3	Santa Cruz
317	4483	Somatostatin Receptor Type 4	Santa Cruz
319	4484	Somatostatin Receptor Type 5	Santa Cruz
321	4552	Tachykinin Receptor 1	Santa Cruz
323	4687	Thrombin Receptor	DPC Biermann/Acris
323	4687	Thrombin Receptor	Research Diagnostics
323	4687	Thrombin Receptor	Santa Cruz
325	4734	Thyrotropin Releasing	Santa Cruz •
	-	Hormone Receptor	All I Divini Tak
327	4944	Angiotensin II Type 1	Alpha Diagnostic Int.
•	-	Receptor	-
327	4944	Angiotensin II Type 1	Biocarta
,		Receptor	
327	4944	Angiotensin II Type: 1	Biogenesis
		Receptor	
327	4944	Angiotensin II Type 1	Capralogics
		Receptor	
327	4944	Angiotensin II Type 1	Chemicon
		Receptor	
327	4944	Angiotensin II Type 1	DPC Biermann/Acris
		Receptor	
327	4944	Angiotensin II Type 1	Fitgerald Industries Int.
		Receptor	
327	4944	Angiotensin II Type 1	Fitzgerald Industries Int.
•		Receptor	
327	4944	Angiotensin II Type 1	Lab Vision Corporation/NeoMarkers
		Receptor	
327	4944	Angiotensin II Type 1	Santa Cruz
		Receptor	
329	4946	Angiotensin II Type 2	Alpha Diagnostic Int.
		Receptor	
329	4946	Angiotensin II Type 2	DPC Biermann/Acris
		Receptor	
329	4946	Angiotensin II Type 2	Santa Cruz
		Receptor	
331	5072	Pyrimidinergic Receptor P2Y4	Chemicon
333	5117	Vasopressin V1A Receptor	Chemicon
335	5118	Vasopressin V1B Receptor	Alpha Diagnostic Int.
335	5118	Vasopressin V1B Receptor	Chemicon
337	5119	Vasopressin V2 Receptor	Alpha Diagnostic Int.
337	5119	Vasopressin V2 Receptor	Chemicon
337	5119	Vasopressin V2 Receptor	Research Diagnostics
347	6031	SIV/HIV Receptor BONZO	Santa Cruz
349	6204	Lysophosphatidic Acid	Exalpha Biologicals
•		Receptor Edg4	
351	6213	C-C Chemokine Receptor 5	Calbiochem
351	6213	C-C Chemokine Receptor 5	Capralogics
351	6213	C-C Chemokine Receptor 5	Chemicon
351	6213	C-C Chemokine Receptor 5	Research Diagnostics
351	6213	C-C Chemokine Receptor 5	Santa Cruz
361	6853	Purinergic Receptor P2Y11	Zymed
		•	

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365	7221	Galanin Receptor GalR2	Alpha Diagnostic Int.
367	7246	Orexin Receptor 1	Alpha Diagnostic Int.
369	7247	Orexin Receptor 2	Alpha Diagnostic Int.
371	8436	Platelet-Activating Factor	Cayman
		Receptor	
371	8436	Platelet-Activating Factor	Santa Cruz
		Receptor	
377	9421	Neuropeptide Y Receptor Type 1	Biogenesis
377	9421	Neuropeptide Y Receptor Type	DPC Biermann/Acris
250	0024		Bassarch Diagnostics
379	9834	Corticotropin releasing factor Receptor 1	Research Diagnostics
379	9834	Corticotropin releasing factor	Santa Cruz
	•	Receptor 1	
385	14198	Interleukin-8 Receptor B	Biosource
385	14198	Interleukin-8 Receptor B	R&D Systems
385	14198	Interleukin-8 Receptor B	Research Diagnostics
385	14198	Interleukin-8 Receptor B	Santa Cruz
387	14641	Calcitonin Receptor	Santa Cruz
389	16041	C-C Chemokine Receptor 6	Research Diagnostics
389	16041	C-C Chemokine Receptor 6	Santa Cruz
391	16599	Smoothened	Research Diagnostics
391	16599	Smoothened	Santa Cruz
397	17535	Gaba(b) Receptor 1	Alpha Diagnostic Int.
397	17535	Gaba(b) Receptor 1	Calbiochem
397	17535	Gaba(b) Receptor 1	Chemicon
397	17535	Gaba(b) Receptor 1	Santa Cruz
423	37498	Xenotropic and Polytropic	Santa Cruz
		Retrovirus Receptor (XPR1)	
435	54053	Gaba(b) Receptor 2	Alpha Diagnostic Int.
435	54053	Gaba(b) Receptor 2	Chemicon
439	56923	Muscarinic acetylcholine Receptor M3	Biogenesis
439	56923		Santa Cruz
432	30,25	Receptor M3	
457	152201	Thyrotropin Receptor	DPC Biermann/Acris
457	152201	Thyrotropin Receptor	Santa Cruz
459	152245	C-C Chemokine Receptor 2	Research Diagnostics
459	152245	C-C Chemokine Receptor 2	Santa Cruz
461	152299	Interleukin-8 Receptor A	Biosource
462	152299		Biosource
461	152299	Interleukin-8 Receptor A	R&D Systems
462	152299	Interleukin-8 Receptor A	R&D Systems
461	152299	Interleukin-8 Receptor A	Research Diagnostics
462	152299	Interleukin-8 Receptor A	Research Diagnostics
461	152299	Interleukin-8 Receptor A	Santa Cruz
462	152299	Interleukin-8 Receptor A	Santa Cruz
468	159973	Vasoactive Intestinal	Exalpha Biologicals
		Polypeptide Receptor 1	
470	160040		Exalpha Biologicals
	·	Polypeptide Receptor 2	
472	160055	Motilin Receptor (GPR38)	Santa Cruz
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	· ·	448/448		•
503	160228	T-Cell Death-Associated Gene 8 (GPR65)	Santa Cruz	
507	160312	Sphingolipid Receptor Edg5		•
515	160329	Proteinase-Activated Receptor 4	Santa Cruz	
535	161214	Galanin Receptor GalR3	Alpha Diagnostic Int.	
537	161221	Urotensin-II Receptor (GPR14)	Santa Cruz	·
546	177168	Cysteinyl Leukotriene CYSLT1 Receptor	Cayman	•
548	177191	Histamine H3 Receptor	Alpha Diagnostic Int.	
548	177191	Histamine H3 Receptor	Chemicon	*
552	180956	Lysophosphatidic Acid Receptor Edg7	Exalpha Biologicals	
562	189900	Sphingolipid Receptor Edg8	Exalpha Biologicals	
628	190774	Histamine H4 Receptor	Alpha Diagnostic Int.	
628	190774	Histamine H4 Receptor	Chemicon	
636	190955	Leukotriene B4 Receptor BLT1	Cayman	

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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

7 A

(54) Title: ANTIGENIC PEPTIDES, SUCH AS FOR G PROTEIN-COUPLED RECEPTORS (GPCRS), ANTIBODIES THERETO, AND SYSTEMS FOR IDENTIFYING SUCH ANTIGENIC PEPTIDES

(57) Abstract: The present invention provides antigenic peptides for GPCRs and antibodies relating thereto, and related systems, methods, compositions, and the like, such as diagnostics and medicaments. Where antibodies against a given GPCR are not known, the present invention provides such antibodies, and preferred antigenic sequences for producing such antibodies. Where antibodies against a given GPCR are known, the present invention provides preferred antigenic peptides for producing antibodies that exhibit improved specificity, affinity or capacity to perform antibody-related actions relative to the known antibodies.

Intel Ial Application No PC I7US 01/50107

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According to	nternational Patent Classification (IPC) or to both national classific	ation and IPC
B. FIELDS	SEARCHED	
IPC 7	cumentation searched (classification system followed by classification CO7K C12N GO1N	
	ion searched other than minimum documentation to the extent that s	
	ata base consulted during the international search (name of data ba	
EMBL,	SEQUENCE SEARCH, EPO-Internal, WPI	Data, BIOSIS, MEDLINE
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C. DOCUME	ENTS CONSIDERED TO BE RELEVANT	
Category °	Citation of document, with indication, where appropriate, of the rel	evant passages Relevant to claim No.
Calogory		
X	ZHOU FENG C ET AL: "Production characterization of an anti-sero receptor antibody which detects 5-HTIA binding sites." MOLECULAR BRAIN RESEARCH, vol. 69, no. 2, 8 June 1999 (199	tonin 1A 15-26 functional
	pages 186-201, XP002222431 ISSN: 0169-328X figure 1; table 1	
		-/
	•	
X Furth	er documents are listed in the continuation of box C.	Patent family members are listed in annex.
° Special cal	legories of cited documents:	"T" later document published after the international filing date
"A" docume	nt defining the general state of the art which is not ered to be of particular relevance	or priority date and not in contact with the application but cited to understand the principle or theory underlying the invention
"E" earlier d	ocument but published on or after the international	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to
filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)		involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the
"O" docume	ent referring to an oral disclosure, use, exhibition or neans	document is combined with one or more other such docu- ments, such combination being obvious to a person skilled
"P" docume	nt published prior to the international filing date but an the priority date claimed	In the art. "&" document member of the same patent family
	actual completion of the international search	Date of mailing of the international search report
6	January 2003	0 8. 04. 2003
Name and n	nailing address of the ISA	Authorized officer
. Taile and II	European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,	Bucka, A
	Fax: (+31-70) 340-3016	Duoisa, ii

Form PGT/ISA/210 (continuation of second sheet) (July 1992)

Inte nal Application No
PU 17 US 01/50107

		PU 17 US 01/50107
C.(Continua	ation) DOCUMENTS CONSIDERED TO BE RELEVANT	
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	RAYMOND JOHN R ET AL: "Immunohistochemical mapping of cellular and subcellular distribution of 5-HT-1A receptors in rat and human kidneys." AMERICAN JOURNAL OF PHYSIOLOGY, vol. 264, no. 1 PART 2, 1993, pages F9-F19, XP001127496 ISSN: 0002-9513 the whole document, in particular figures	1-10, 15-26
	1, 3	
Y	VERDOT L ET AL: "PRODUCTION OF ANTI-PEPTIDE ANTIBODIES DIRECTED AGAINST THE FIRST AND THE SECOND EXTRACELLULAR LOOP OF THE HUMAN SEROTONIN 5-HT1A RECEPTOR"	1-10, 15-26
	BIOCHIMIE, MASSON, PARIS, FR, vol. 76, no. 1, 1994, pages 165-170, XP008009332 ISSN: 0300-9084 the whole document	
γ	TODD E ANTHONY AND EFRAIAN C AZMITIA: "Molecular characterization of antipeptide antibodies against the 5-HT1A receptor: Evidence for state-dependent antibody binding."	1-10, 15-26
	MOLECULAR BRAIN RESEARCH, vol. 50, no. 1-2, 15 October 1997 (1997-10-15), pages 277-284, XP002222432 ISSN: 0169-328X the whole document	
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	the whole document BACKSTROM JON R ET AL: "Generation of anti-peptide antibodies against serotonin 5-HT2A and 5-HT2C receptors."	1-10, 15-26
	JOURNAL OF NEUROSCIENCE METHODS, vol. 77, no. 1, 7 November 1997 (1997-11-07), pages 109-117, XP002222433 ISSN: 0165-0270 the whole document	
	-/- -	

Inte al Application No PC17US 01/50107

.(Continua ategory °	tion) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages		Relevant to claim No.
	EASON MARGARET G ET AL: "Identification of a G-s coupling domain in the amino terminus of the third intracellular loop of the alpha-2A-adrenergic receptor: Evidence for distinct structural determinants that confer G-s versus G-i		1-10, 15-26
	coupling." JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 270, no. 42, 1995, pages 24753-24760, XP002222434 ISSN: 0021-9258 the whole document		
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national application No. PCT/US 01/50107

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)	
This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:	-
1. X Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:	
Although claims 19 and 20 are directed to a diagnostic method practised on the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.	he
2. Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:	
3. Claims Nos.:	•
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).	
Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)	
This International Searching Authority found multiple inventions in this international application, as follows:	
a diameter application, as latitudes.	
see additional sheet	
1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.	
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.	
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:	
4. No required additional search fees were timely paid by the applicant. Consequently, this international Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:	-
1-10, 15-26 (all partially)	
Remark on Protest The additional search fees were accompanied by the applicant's protest.	
No protest accompanied the payment of additional search fees.	
	1

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

Invention 1: claims 1-10, 15-26, all partially

an isolated antigenic peptide having the amino acid sequence SEQ ID NO: 692, nucleic acids encoding said peptide, antibodies directed against said peptide, kits containing said antibodies

Inventions 2 to 1600: claims 1-26, all partially and in so far as applicable

each separate, individual invention relates to an isolated antigenic peptide, nucleic acids encoding said peptide, antibodies directed against said peptide, kits containing said antibodies, wherein invention 2 is represented by the peptide having the amino acid sequence SEQ ID NO: 693, invention 3 is represented by the peptide having the amino acid sequence SEQ ID NO: 694, continuing to invention 1600, which is represented by the peptide having the amino acid sequence SEQ ID NO: 2292

Invention 1601: claims 27-66

a method of identifying an amino acid sequence of an antigenic peptide derived from a candidate polypeptide, peptides identified by that method, antibodies directed against said peptides